



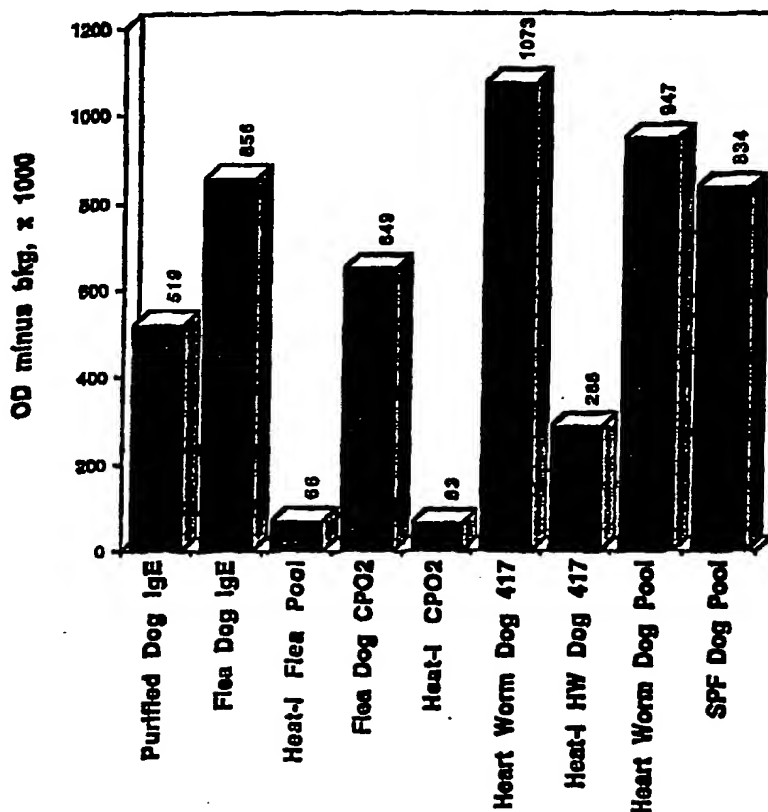
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(54) Title: METHOD TO DETECT IgE

(57) Abstract

The present invention includes a method to detect IgE using a human Fc epsilon receptor (FcεR) to detect IgE antibodies in a biological sample from a cat, a dog, or a horse. The present invention also relates to kits to perform such methods.



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METHOD TO DETECT IgE

Field of the Invention

The present invention relates to a novel method to detect epsilon immunoglobulin (IgE). The present invention also includes novel kits to detect IgE as well as methods to produce the detection reagent.

Background of the Invention

Diagnosis of disease and determination of treatment efficacy are important tools in medicine. In particular, detection of IgE production in an animal can be indicative of disease. Such diseases include, for example, allergy, atopic disease, hyper IgE syndrome, internal parasite infections and B cell neoplasia. In addition, detection of IgE production in an animal following a treatment is indicative of the efficacy of the treatment, such as when using treatments intended to disrupt IgE production.

Until the discovery of the present invention, detection of IgE in samples obtained from non-human animals has been hindered by the absence of suitable reagents for detection of IgE. Various compounds have been used to detect IgE in IgE-containing compositions. In particular, antibodies that bind selectively to epsilon idiotype antibodies (i.e., anti-IgE antibodies) have been used to detect IgE. These anti-IgE antibodies, however, can cross-react with other antibody idiotypes, such as gamma isotype antibodies. The discovery of the present invention includes the use of a Fc epsilon receptor (Fc ϵ R) molecule to detect the presence of IgE in a putative IgE-containing composition. A Fc ϵ R molecule provides an advantage over, for example anti-IgE antibodies, to detect IgE because a Fc ϵ R molecule can bind to an IgE with more specificity (i.e., less idiotype cross-reactivity) and more sensitivity (i.e., affinity) than anti-IgE binding antibodies.

Lowenthal et al., 1993, *Annals of Allergy* 71:481-484, dog serum can transfer cutaneous reactivity to a human. While it is possible that Lowenthal et al. properly teach the binding of human Fc ϵ R to canine IgE. Lowenthal et al., however, do not provide data defining the particular cellular proteins responsible for the transfer of cutaneous reactivity. As such, a skilled artisan would conclude that the transfer of cutaneous reactivity taught by Lowenthal et al. could be due to a variety of different molecular interactions and that the conclusion drawn by Lowenthal et al. is merely an

interpretation. In addition, Lowenthal et al. do not teach the use of purified human Fc_εR to detect canine IgE. The subunits of human Fc_εR have been known as early as 1988 and have never been used to detect canine, feline or equine IgE. Indeed, U.S. Patent No. 4,962,035, to Leder et al., issued October 9, 1990, discloses human Fc_εR but does not
5 disclose the use of such a human Fc_εR to detect human or non-human IgE. The use of purified human Fc_εR avoids complications presented by use of Fc_εR bound to a cell, such as non-specific binding of the Fc_εR-bearing cell due to additional molecules present on the cell membrane. That purified human Fc_εR detects non-human IgE is unexpected because inter-species binding between a Fc_εR and an IgE is not predictable. For
10 example, human Fc_εR binds to rat IgE but rat Fc_εR does not bind to human IgE.

The high affinity Fc_εR consists of three protein chains, alpha, beta and gamma. Prior investigators have disclosed the nucleic acid sequence for: the alpha chain (Kochan et al., *Nucleic Acids Res.* 16:3584, 1988; Shimizu et al., *Proc. Natl. Acad. Sci. USA* 85:1907-1911, 1988; and Pang et al., *J. Immunol.* 151:6166-6174, 1993); the beta chain
15 (Kuster et al., *J. Biol. Chem.* 267:12782-12787, 1992); and the gamma chain (Kuster et al., *J. Biol. Chem.* 265:6448-6452, 1990).

Thus, methods and kits are needed in the art that will provide specific detection of non-human IgE.

Summary of the Invention

20 The present invention includes detection methods and kits that detect IgE. One embodiment of the present invention is a method to detect IgE comprising: (a) contacting an isolated human Fc_ε receptor (Fc_εR) molecule with a putative IgE-containing composition under conditions suitable for formation of a Fc_εR molecule:IgE complex, wherein the IgE is selected from the group consisting of canine IgE, feline IgE
25 and equine IgE; and (b) determining the presence of IgE by detecting the Fc_εR molecule:IgE complex, the presence of the Fc_εR molecule:IgE complex indicating the presence of IgE. A preferred Fc_εR molecule in which a carbohydrate group of the Fc_εR molecule is conjugated to biotin.

Another embodiment of the present invention is a method to detect IgE
30 comprising: (a) contacting a recombinant cell with a putative IgE-containing composition under conditions suitable for formation of a recombinant cell:IgE complex,

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in which the recombinant cell includes: a recombinant cell expressing a human $Fc_\epsilon R$ molecule; and a recombinant cell expressing an antibody that binds selectively to an IgE including canine IgE, feline IgE and equine IgE; and (b) determining the presence of IgE by detecting the recombinant cell:IgE complex, the presence of the recombinant cell:IgE complex indicating the presence of IgE. A preferred recombinant cell includes a RBL-h $Fc_\epsilon R$ cell.

Another embodiment of the present invention is a method to detect flea allergy dermatitis comprising: (a) immobilizing a flea allergen on a substrate; (b) contacting the flea allergen with a putative IgE-containing composition under conditions suitable for formation of an antigen:IgE complex bound to said substrate; (c) removing non-bound material from the substrate under conditions that retain antigen:IgE complex binding to the substrate; and (c) detecting the presence of the antigen:IgE complex by contacting the antigen:IgE complex with a $Fc_\epsilon R$ molecule. Preferably, the flea allergen is a flea saliva antigen and more preferably flea saliva products and/or flea saliva proteins.

The present invention also includes a kit for performing methods of the present invention. One embodiment is a kit for detecting IgE comprising a human Fc_ϵ receptor ($Fc_\epsilon R$) molecule and a means for detecting an IgE including canine IgE, feline IgE and equine IgE. Another embodiment is a general allergen kit comprising an allergen common to all regions of the United States and a human Fc_ϵ receptor ($Fc_\epsilon R$) molecule. Another embodiment is a kit for detecting flea allergy dermatitis comprising a human Fc_ϵ receptor ($Fc_\epsilon R$) molecule and a flea allergen.

Another embodiment of the present invention is an isolated human Fc_ϵ receptor ($Fc_\epsilon R$) alpha chain protein, in which a carbohydrate group of the $Fc_\epsilon R$ alpha chain protein is conjugated to biotin. A preferred $Fc_\epsilon R$ alpha chain protein comprises Ph $Fc_\epsilon R\alpha_{172}$ -BIOT.

Brief Description of the Figures

Fig. 1 depicts ELISA results using biotinylated alpha chain of human $Fc_\epsilon R$ to detect canine IgE antibodies.

Fig. 2 depicts ELISA results using biotinylated alpha chain of human $Fc_\epsilon R$ to detect plant allergen-specific canine IgE antibodies.

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Fig. 3 depicts ELISA results using biotinylated alpha chain of human $Fc_\epsilon R$ to detect human or canine IgE antibodies.

Fig. 4 depicts ELISA results using biotinylated alpha chain of human $Fc_\epsilon R$ to detect flea allergen-specific canine IgE antibodies.

5 Fig. 5 depicts ELISA results using biotinylated alpha chain of human $Fc_\epsilon R$ to detect flea allergen-specific and heartworm antigen-specific canine IgE antibodies.

Fig. 6 depicts ELISA results using biotinylated alpha chain of human $Fc_\epsilon R$ to detect flea saliva-specific canine IgE antibodies.

10 Fig. 7 depicts ELISA results using biotinylated alpha chain of human $Fc_\epsilon R$ to detect heartworm antigen-specific feline IgE antibodies.

Fig. 8 depicts ELISA results using biotinylated alpha chain of human $Fc_\epsilon R$ to detect heartworm antigen-specific feline IgE antibodies.

Fig. 9 depicts ELISA results using biotinylated alpha chain of human $Fc_\epsilon R$ to detect antigen-specific equine IgE antibodies.

15 Fig. 10 depicts ELISA results using basophilic leukemia cells expressing alpha chain of human $Fc_\epsilon R$ to detect canine IgE antibodies in sera from heartworm-infected dogs.

20 Fig. 11 depicts ELISA results using basophilic leukemia cells expressing alpha chain of human $Fc_\epsilon R$ to detect canine IgE antibodies in sera from flea saliva sensitized dogs.

Detailed Description of the Invention

The present invention relates to the discovery that purified high affinity human Fc epsilon receptor (i.e., $Fc_\epsilon RI$; referred to herein as $Fc_\epsilon R$) can be used in certain non-human (i.e., canine, feline or equine) epsilon-immunoglobulin (referred to herein as IgE or IgE antibody)-based detection (e.g., diagnostic, screening) methods and kits. The use of human $Fc_\epsilon R$ to detect non-human IgE is unexpected because canine, feline and equine immune systems are different from the human immune system, as well as from each other (i.e., molecules important to the immune system usually are species specific).

30 One embodiment of the present invention is a method to detect a non-human IgE using an isolated human $Fc_\epsilon R$ molecule. It is to be noted that the term "a" entity or "an" entity refers to one or more of that entity; for example, a protein refers to one or more

proteins or at least one protein. As such, the terms "a" (or "an"), "one or more" and "at least one" can be used interchangeably herein. It is also to be noted that the terms "comprising", "including", and "having" can be used interchangeably. It is also to be noted that the terms "comprising", "including", and "having" can be used interchangeably. Furthermore, a compound "selected from the group consisting of" refers to one or more of the compounds in the list that follows, including mixtures (i.e., combinations) of two or more of the compounds.

According to the present invention, an isolated, or biologically pure, $Fc_\epsilon R$ molecule, is a molecule that has been removed from its natural milieu. As such, "isolated" and "biologically pure" do not necessarily reflect the extent to which the molecule has been purified. An isolated human $Fc_\epsilon R$ molecule of the present invention can be obtained from its natural source (e.g., from a human mast cell), can be produced using recombinant DNA technology or can be produced by chemical synthesis.

A $Fc_\epsilon R$ molecule (also referred to herein as $Fc_\epsilon R$ or $Fc_\epsilon R$ protein) of the present invention can be a full-length protein, a portion of a full-length protein or any homolog of such a protein. As used herein, a protein can be a polypeptide or a peptide. A $Fc_\epsilon R$ molecule of the present invention can comprise a complete $Fc_\epsilon R$ (i.e., alpha, beta and gamma $Fc_\epsilon R$ chains), an alpha $Fc_\epsilon R$ chain (also referred to herein as $Fc_\epsilon R$ α chain) or portions thereof. Preferably, a $Fc_\epsilon R$ molecule comprises at least a portion of a $Fc_\epsilon R$ α chain that binds to IgE, i.e., that is capable of forming an immunocomplex with an IgE constant region. Preferably, a $Fc_\epsilon R$ molecule of the present invention binds to IgE with an affinity of about $K_A \approx 10^8$, more preferably with an affinity of about $K_A \approx 10^9$ and even more preferably with an affinity of about $K_A \approx 10^{10}$.

An isolated $Fc_\epsilon R$ molecule of the present invention, including a homolog, can be identified in a straight-forward manner by the $Fc_\epsilon R$ molecule's ability to form an immunocomplex with an IgE. Examples of $Fc_\epsilon R$ homologs include $Fc_\epsilon R$ proteins in which amino acids have been deleted (e.g., a truncated version of the protein, such as a peptide), inserted, inverted, substituted and/or derivatized (e.g., by glycosylation, phosphorylation, acetylation, myristoylation, prenylation, palmitoylation, amidation and/or addition of glycerophosphatidyl inositol) such that the homolog includes at least one epitope capable of forming an immunocomplex with an IgE.

Fc_εR homologs can be the result of natural allelic variation or natural mutation. Fc_εR homologs of the present invention can also be produced using techniques known in the art including, but not limited to, direct modifications to the protein or modifications to the gene encoding the protein using, for example, classic or recombinant DNA techniques to effect random or targeted mutagenesis.

According to the present invention, a human Fc_εR α chain of the present invention is encoded by at least a portion of the nucleic acid sequence of the coding strand of a cDNA encoding a full-length Fc_εR α chain protein represented herein as SEQ ID NO:1, the portion at least encoding the IgE binding site of the Fc_εR α chain protein.

10 The double-stranded nucleic acid molecule including both the coding strand having SEQ ID NO:1 and the complementary non-coding strand (the nucleic acid sequence of which can be readily determined by one skilled in the art and is shown herein as SEQ ID NO:3) is referred to herein as Fc_εR nucleic acid molecule nhFc_εRα₁₁₉₈. Translation of SEQ ID NO:1 suggests that nucleic acid molecule nhFc_εRα₁₁₉₈ encodes a full-length Fc_εR α

15 chain protein of about 257 amino acids, referred to herein as PhFc_εRα₂₅₇, represented by SEQ ID NO:2, assuming an open reading frame having an initiation (start) codon spanning from nucleotide 107 through nucleotide 109 of SEQ ID NO:1 and a termination (stop) codon spanning from nucleotide 878 through nucleotide 880 of SEQ ID NO:1. The coding region encoding PhFc_εRα₂₅₇, including the stop codon, is

20 represented by nucleic acid molecule nhFc_εRα₇₇₄, having a coding strand with the nucleic acid sequence represented herein as SEQ ID NO:4. The complement of SEQ ID NO:4 is represented herein as SEQ ID NO:5. SEQ ID NO:1 encodes a signal peptide of about 25 amino acids as well as a mature protein of about 232 amino acids, denoted herein as PhFc_εRα₂₃₂, the amino acid sequence of which is represented herein as SEQ ID

25 NO:6. The nucleic acid molecule encoding the apparent mature protein is referred to as nhFc_εRα₆₉₉, the nucleic acid sequence of the coding strand of which is denoted herein as SEQ ID NO:7. SEQ ID NO:1 also encodes a hydrophobic transmembrane domain and a cytoplasmic tail which as a group extend from amino acid 205 to amino acid 257 of SEQ ID NO:2. Knowledge of these nucleic acid and amino acid sequences allows one skilled

30 in the art to make modifications to the respective nucleic acid molecules and proteins to, for example, develop a Fc_εR α chain protein with increased solubility and/or a truncated

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protein (e.g., a peptide) capable of detecting IgE, e.g., $\text{PhFc}_\epsilon\text{R}\alpha_{197}$ and $\text{PhFc}_\epsilon\text{R}\alpha_{172}$. Preferred $\text{Fc}_\epsilon\text{R}$ molecules include $\text{PhFc}_\epsilon\text{R}\alpha_{257}$, $\text{PhFc}_\epsilon\text{R}\alpha_{197}$, $\text{PhFc}_\epsilon\text{R}\alpha_{232}$ and $\text{PhFc}_\epsilon\text{R}\alpha_{172}$. Preferred nucleic acid molecules to encode a $\text{Fc}_\epsilon\text{R}$ molecules include $\text{nhFc}_\epsilon\text{R}\alpha_{774}$, $\text{nhFc}_\epsilon\text{R}\alpha_{1198}$, $\text{nhFc}_\epsilon\text{R}\alpha_{612}$, $\text{nhFc}_\epsilon\text{R}\alpha_{591}$, $\text{nhFc}_\epsilon\text{R}\alpha_{699}$ and/or $\text{nhFc}_\epsilon\text{R}\alpha_{516}$.

- 5 Isolated $\text{Fc}_\epsilon\text{R}$ molecule protein of the present invention can be produced by culturing a cell capable of expressing the protein under conditions effective to produce the protein, and recovering the protein. A preferred cell to culture is a recombinant cell that is capable of expressing the protein, the recombinant cell being produced by transforming a host cell with one or more nucleic acid molecules of the present
- 10 invention. Transformation of a nucleic acid molecule into a cell can be accomplished by any method by which a nucleic acid molecule can be inserted into the cell. Transformation techniques include, but are not limited to, transfection, electroporation, microinjection, lipofection, adsorption, and protoplast fusion. A recombinant cell may remain unicellular or may grow into a tissue, organ or a multicellular organism.
- 15 Transformed nucleic acid molecules of the present invention can remain extrachromosomal or can integrate into one or more sites within a chromosome of the transformed (i.e., recombinant) cell in such a manner that their ability to be expressed is retained. Suitable and preferred nucleic acid molecules with which to transform a cell are as disclosed herein for suitable and preferred $\text{Fc}_\epsilon\text{R}$ nucleic acid molecules per se.
- 20 Particularly preferred nucleic acid molecules to include in recombinant cells of the present invention include $\text{nhFc}_\epsilon\text{R}\alpha_{774}$, $\text{nhFc}_\epsilon\text{R}\alpha_{1198}$, $\text{nhFc}_\epsilon\text{R}\alpha_{612}$, $\text{nhFc}_\epsilon\text{R}\alpha_{591}$, $\text{nhFc}_\epsilon\text{R}\alpha_{699}$ and/or $\text{nhFc}_\epsilon\text{R}\alpha_{516}$.

- Suitable host cells to transform include any cell that can be transformed with a nucleic acid molecule of the present invention. Host cells can be either untransformed
- 25 cells or cells that are already transformed with at least one nucleic acid molecule. Host cells of the present invention either can be endogenously (i.e., naturally) capable of producing a $\text{Fc}_\epsilon\text{R}$ molecule protein of the present invention or can be capable of producing such proteins after being transformed with at least one nucleic acid molecule of the present invention. Host cells of the present invention can be any cell capable of
- 30 producing at least one protein of the present invention, and include bacterial, fungal

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(including yeast), parasite (including protozoa and ectoparasite), insect, other animal and plant cells.

Preferably, a recombinant cell is transfected with a recombinant molecule of the present invention is a molecule that can include at least one of any nucleic acid molecule heretofore described operatively linked to at least one of any transcription control sequence capable of effectively regulating expression of the nucleic acid molecule(s) in the cell to be transformed, examples of which are disclosed herein. A particularly preferred recombinant molecule includes pVL-nhFc_εRα₆₁₂. Details regarding the production of Fc_εR molecule nucleic acid molecule-containing recombinant molecules are disclosed herein. Particularly preferred recombinant cell of the present invention includes *Trichoplusia ni*-pVL-nhFc_εRα₆₁₂.

A Fc_εR molecule of the present invention can include chimeric molecules comprising a portion of a Fc_εR molecule that binds to an IgE and a second molecule that enables the chimeric molecule to be bound to a substrate in such a manner that the Fc_εR portion binds to IgE in essentially the same manner as a Fc_εR molecule that is not bound to a substrate. An example of a suitable second molecule includes a portion of an immunoglobulin molecule.

A Fc_εR molecule of the present invention can be contained in a formulation, herein referred to as a Fc_εR formulation. For example, a Fc_εR can be combined with a buffer in which the Fc_εR is solubilized, and/or a carrier. Suitable buffers and carriers are known to those skilled in the art. Examples of suitable buffers include any buffer in which a Fc_εR can function to selectively bind to IgE, such as, but not limited to, phosphate buffered saline, water, saline, phosphate buffer, bicarbonate buffer, HEPES buffer (N-2-hydroxyethylpiperazine-N'-2-ethanesulfonic acid buffered saline), TES buffer (Tris-EDTA buffered saline), Tris buffer and TAE buffer (Tris-acetate-EDTA). Examples of carriers include, but are not limited to, polymeric matrices, toxoids, and serum albumins, such as bovine serum albumin. Carriers can be in mixed with Fc_εR or conjugated (i.e., attached) to Fc_εR in such a manner as to not substantially interfere with the ability of the Fc_εR to selectively bind to IgE.

A Fc_εR of the present invention can be bound to the surface of a cell expressing the Fc_εR. A preferred Fc_εR-bearing cell includes a recombinant cell expressing a nucleic

acid molecule encoding a human $\text{Fc}_\epsilon\text{R}$ alpha chain of the present invention. A more preferred recombinant cell of the present invention expresses a nucleic acid molecule that encodes at least one of the following proteins: $\text{PhFc}_\epsilon\text{R}\alpha_{257}$ and $\text{PhFc}_\epsilon\text{R}\alpha_{232}$. An even more preferred recombinant cell expresses a nucleic acid molecule including $\text{nhFc}_\epsilon\text{R}\alpha_{612}$,
5 $\text{nhFc}_\epsilon\text{R}\alpha_{591}$, $\text{nhFc}_\epsilon\text{R}\alpha_{699}$ and/or $\text{nhFc}_\epsilon\text{R}\alpha_{516}$ with a recombinant cell expressing a nucleic acid molecule comprising a nucleic acid sequence including SEQ ID NO:1 or SEQ ID NO:4, or a nucleic acid molecule comprising an allelic variant of a nucleic acid molecule comprising SEQ ID NO:1 or SEQ ID NO:4, being even more preferred. An even more preferred recombinant cell is a RBL-h $\text{Fc}_\epsilon\text{R}$ cell.

10 In addition, a $\text{Fc}_\epsilon\text{R}$ formulation of the present invention can include not only a $\text{Fc}_\epsilon\text{R}$ but also one or more additional antigens or antibodies useful in detecting IgE. As used herein, an antigen refers to any molecule capable of being selectively bound by an antibody. As used herein, specific binding of a first molecule to a second molecule refers to the ability of the first molecule to preferentially bind (e.g., having higher
15 affinity higher avidity) to the second molecule when compared to the ability of a first molecule to bind to a third molecule. The first molecule need not necessarily be the natural ligand of the second molecule. Examples of such antibodies include, but are not limited to, antibodies that bind selectively to the constant region of an IgE heavy (i.e., anti-IgE isotype antibody) or antibodies that bind selectively to an IgE having a specific
20 antigen specificity (i.e., anti-IgE idiotypic antibody). Examples of such antigens include any antigen known to induce the production of IgE. Preferred antigens include allergens and parasite antigens. Allergens of the present invention are preferably derived from fungi, trees, weeds, shrubs, grasses, wheat, corn, soybeans, rice, eggs, milk, cheese, bovines (or cattle), poultry, swine, sheep, yeast, fleas, flies, mosquitos, mites, midges,
25 biting gnats, lice, bees, wasps, ants, true bugs or ticks. A suitable flea allergen includes an allergen derived from a flea, in particular flea saliva antigen. A preferred flea allergen includes a flea saliva antigen Preferred flea saliva antigens include antigens such as those disclosed in PCT Patent Publication No. WO 96/11271, published April 18, 1996, by Frank et al. (this publication is incorporated by reference herein in its
30 entirety), with flea saliva products and flea saliva proteins being particularly preferred. According to the present invention, a flea saliva protein includes a protein produced by

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recombinant DNA methods, as well as proteins isolated by other methods disclosed in PCT Patent Publication No. WO 96/11271.

- Preferred general allergens include those derived from grass, Meadow Fescue, Curly Dock, plantain, Mexican Firebush, Lamb's Quarters, pigweed, ragweed, sage, elm, 5 cocklebur, Box Elder, walnut, cottonwood, ash, birch, cedar, oak, mulberry, cockroach, *Dermataphagoides*, *Alternaria*, *Aspergillus*, *Cladosporium*, *Fusarium*, *Helminthosporium*, *Mucor*, *Penicillium*, *Pullularia*, *Rhizopus* and/or *Tricophyton*. More preferred general allergens include those derived from Johnson Grass, Kentucky Blue Grass, Meadow Fescue, Orchard Grass, Perennial Rye Grass, Redtop Grass, Timothy 10 Grass, Bermuda Grass, Brome Grass, Curly Dock, English Plantain, Mexican Firebush, Lamb's Quarters, Rough Pigweed Short Ragweed, Wormwood Sage, American Elm, Common Cocklebur, Box Elder, Black Walnut, Eastern Cottonwood, Green Ash, River Birch, Red Cedar, Red Oak, Red Mulberry, Cockroach, *Dermataphagoides farinae*, *Alternaria alternata*, *Aspergillus fumigatus*, *Cladosporium herbarum*, *Fusarium* 15 *vasinfectum*, *Helminthosporium sativum*, *Mucor recemosus*, *Penicillium notatum*, *Pullularia pullulans*, *Rhizopus nigricans* and/or *Tricophyton* spp. Preferred tropical allergens include those derived from Bermuda Grass, June Bluegrass, Annual Bluegrass, Orchard Grass, Perennial Rye Grass, Timothy Grass, Meadow Fescue, Common Cocklebur, Yellow Dock, Sheep Sorrel, English Plantain, Lamb's Quarters, Rough 20 Pigweed, Russian Thistle, Short Ragweed, Red Cedar, Cat Epithelium, Arizona Cypress, Bald Cypress, Date Palm, Australian Pine, Eucalyptus, Mango, Acacia, Grama Grass, Nettle, Western Cottonwood, Saltgrass, *Dermataphagoides pteronyssinus*, *Aureobasidium pullans*, *Penicillium notatum*, *Penicillium chrysogenum*, *Drechslera sorokiniana*, *Fusarium roseum*, *Cladosporium sphaerospermum*, *Aspergillus fumigatus*, 25 *Alternaria tenuis* *Dermataphagoides farinae* and *Stemphyllium sarciniforme*. Preferred desert allergens include those derived from Bahia Grass, Smooth Brome, Johnson Grass, Redtop Grass, False Ragweed, Carelessweed, Greasewood, Rough Marsh Elder, Kochia, Tall Ragweed, Western Ragweed, Slender Ragweed, Common Sage, Prairie Sage, Mugwort Sage and Shadscale. Preferred parasite antigens include, but are not limited to, 30 helminth antigens, in particular heartworm antigens, such as Di33 (described in U.S. Patent Application Serial No. 08/715,628, filed September 18, 1996, to Grieve et al.).

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The term "derived from" refers to a natural allergen of such plants or organisms (i.e., an allergen directly isolated from such plants or organisms), as well as, non-natural allergens of such plants or organisms that possess at least one epitope capable of eliciting an immune response against an allergen (e.g., produced using recombinant DNA technology or by chemical synthesis).

The present invention also includes human $Fc_\epsilon R$ mimetopes and use thereof to detect IgE. In accordance with the present invention, a "mimotope" refers to any compound that is able to mimic the ability of a $Fc_\epsilon R$ molecule to bind to IgE. A mimotope can be a peptide that has been modified to decrease its susceptibility to degradation but that still retains IgE-binding activity. Other examples of mimetopes include, but are not limited to, carbohydrate-based compounds, lipid-based compounds, nucleic acid-based compounds, natural organic compounds, synthetically derived organic compounds, anti-idiotypic antibodies and/or catalytic antibodies, or fragments thereof. A mimotope can be obtained by, for example, screening libraries of synthetic compounds for compounds capable of binding to IgE. A mimotope can also be obtained by, for example, rational drug design. In a rational drug design procedure, the three-dimensional structure of a compound of the present invention can be analyzed by, for example, nuclear magnetic resonance (NMR) or x-ray crystallography. The three-dimensional structure can then be used to predict structures of potential mimetopes by, for example, computer modeling. The predicted mimotope structures can then be produced by, for example, chemical synthesis, recombinant DNA technology, or by isolating a mimotope from a natural source. Specific examples of $Fc_\epsilon R$ mimetopes include anti-idiotypic antibodies, oligonucleotides produced using Selex technology, peptides identified by random screening of peptide libraries and proteins identified by phage display technology.

One embodiment of the present invention is a method to detect non-human IgE which includes the steps of: (a) contacting an isolated human $Fc_\epsilon R$ receptor ($Fc_\epsilon R$) molecule with a putative IgE-containing composition under conditions suitable for formation of an $Fc_\epsilon R$ molecule:IgE complex; and (b) determining levels of IgE by detecting said $Fc_\epsilon R$ molecule:IgE complex. Presence of such a $Fc_\epsilon R$ molecule:IgE complex indicates that the animal is producing IgE. Preferred non-human IgE to detect

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using a human $\text{Fc}_\epsilon\text{R}$ molecule include canine IgE, feline IgE and equine IgE. The present method can further include the step of determining whether an IgE complexed with a $\text{Fc}_\epsilon\text{R}$ molecule is heat labile. Methods to determine heat lability of IgE are disclosed in the Examples section. Preferably, an IgE is heat labile when incubated at about 56°C for about 4 hours. Without being bound by theory, Applicants believe that heat labile forms of IgE bind to certain allergens and non-heat labile forms of IgE bind to other types of allergens. As such, detection of heat labile IgE compared with non-heat labile IgE can be used to discriminate between allergen sensitivities. For example, Applicants believe that IgE antibodies that bind to certain flea allergens and heartworm allergens are heat labile while IgE antibodies that bind to certain plant allergens are not heat labile. Thus, the presence of non-heat labile IgE can indicate that an animal is sensitive to certain plant allergens but not to certain flea or heartworm allergens. Moreover, Applicants believe that identification of heat labile IgE and non-heat labile IgE using a human $\text{Fc}_\epsilon\text{R}$ suggests the presence of different sub-populations of IgE that may or may not have substantially similar structures to known IgE. As such, a $\text{Fc}_\epsilon\text{R}$ molecule of the present invention may be useful for detecting molecules bound by the $\text{Fc}_\epsilon\text{R}$ molecule but not identical to a known IgE.

As used herein, canine refers to any member of the dog family, including domestic dogs, wild dogs and zoo dogs. Examples of dogs include, but are not limited to, domestic dogs, wild dogs, foxes, wolves, jackals and coyotes. As used herein, a feline refers to any member of the cat family, including domestic cats, wild cats and zoo cats. Examples of cats include, but are not limited to, domestic cats, lions, tigers, leopards, panthers, cougars, bobcats, lynx, jaguars, cheetahs, and servals. As used herein, equine refers to any member of the horse family, including horses, donkeys, mules and zebras.

As used herein, the term "contacting" refers to combining or mixing, in this case a putative IgE-containing composition with a human $\text{Fc}_\epsilon\text{R}$ molecule. For formation of a complex between a $\text{Fc}_\epsilon\text{R}$ and an IgE refers to the ability of the $\text{Fc}_\epsilon\text{R}$ to selectively bind to the IgE in order to form a stable complex that can be measured (i.e., detected). As used herein, the term selectively binds to an IgE refers to the ability of a $\text{Fc}_\epsilon\text{R}$ of the present invention to preferentially bind to IgE, without being able to substantially bind to

other antibody isotypes. Binding between a Fc_εR and an IgE is effected under conditions suitable to form a complex; such conditions (e.g., appropriate concentrations, buffers, temperatures, reaction times) as well as methods to optimize such conditions are known to those skilled in the art, and examples are disclosed herein. Examples of complex
5 formation conditions are also disclosed in, for example, in Sambrook et al., *Molecular Cloning: A Laboratory Manual*, Cold Spring Harbor Labs Press, 1989, the reference Sambrook et al., *ibid.*, is incorporated by reference herein in its entirety.

As used herein, the term "detecting complex formation" refers to determining if any complex is formed, i.e., assaying for the presence (i.e., existence) of a complex. If
10 complexes are formed, the amount of complexes formed can, but need not be, determined. Complex formation, or selective binding, between Fc_εR and any IgE in the composition can be measured (i.e., detected, determined) using a variety of methods standard in the art (see, for example, Sambrook et al. *ibid.*), examples of which are disclosed herein.

15 In one embodiment, a putative IgE-containing composition of the present method includes a biological sample from an animal. A suitable biological sample includes, but is not limited to, a bodily fluid composition or a cellular composition. A bodily fluid refers to any fluid that can be collected (i.e., obtained) from an animal, examples of which include, but are not limited to, blood, serum, plasma, urine, tears, aqueous humor,
20 central nervous system fluid (CSF), saliva, lymph, nasal secretions, milk and feces. Such a composition of the present method can, but need not be, pretreated to remove at least some of the non-IgE isotypes of immunoglobulin and/or other proteins, such as albumin, present in the fluid. Such removal can include, but is not limited to, contacting the bodily fluid with a material, such as Protein G, to remove IgG antibodies and/or
25 affinity purifying IgE antibodies from other components of the body fluid by exposing the fluid to, for example, Concanavalin A. In another embodiment, a composition includes collected bodily fluid that is pretreated to concentrate immunoglobulin contained in the fluid. For example, immunoglobulin contained in a bodily fluid can be precipitated from other proteins using ammonium sulfate. A preferred composition of
30 the present method is serum.

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In another embodiment, a composition of the present method includes an IgE-producing cell. Such a cell can have IgE bound to the surface of the cell and/or can secrete IgE. Examples of such cells include basophil cells and myeloma cells. IgE can be bound to the surface of a cell either directly to the membrane of a cell or bound to a molecule (e.g., an antigen) bound to the surface of the cell.

A complex can be detected in a variety of ways including, but not limited to use of one or more of the following assays: an enzyme-linked immunoassay, a radioimmunoassay, a fluorescence immunoassay, a chemiluminescent assay, a lateral flow assay, an agglutination assay, a particulate-based assay (e.g., using particulates such as, but not limited to, magnetic particles or plastic polymers, such as latex or polystyrene beads), an immunoprecipitation assay, a BioCore™ assay (e.g., using colloidal gold) and an immunoblotting assay (e.g., a western blot). Such assays are well known to those skilled in the art. Assays can be used to give qualitative or quantitative results depending on how they are used. Some assays, such as agglutination, particulate separation, and immunoprecipitation, can be observed visually (e.g., either by eye or by a machine, such as a densitometer or spectrophotometer) without the need for a detectable marker. In other assays, conjugation (i.e., attachment) of a detectable marker to the Fc_εR or to a reagent that selectively binds to the Fc_εR or to the IgE being detected (described in more detail below) aids in detecting complex formation. Examples of detectable markers include, but are not limited to, a radioactive label, a fluorescent label, a chemiluminescent label, a chromophoric label or a ligand. A ligand refers to a molecule that binds selectively to another molecule. Preferred detectable markers include, but are not limited to, fluorescein, a radioisotope, a phosphatase (e.g., alkaline phosphatase), biotin, avidin, a peroxidase (e.g., horseradish peroxidase) and biotin-related compounds or avidin-related compounds (e.g., streptavidin or ImmunoPure® NeutrAvidin). Preferably, biotin is conjugated to an alpha chain of a Fc_εR. Preferably a carbohydrate group of the Fc_εR alpha chain is conjugated to biotin. A preferred Fc_εR molecule conjugated to biotin comprises PhFc_εRα₁₇₂-BIOT (the production of which is described in the Examples section).

In one embodiment, a complex is detected by contacting a putative IgE-containing composition with a Fc_εR molecule that is conjugated to a detectable marker.

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A suitable detectable marker to conjugate to a $Fc_\epsilon R$ molecule includes, but is not limited to, a radioactive label, a fluorescent label, a chemiluminescent label or a chromophoric label. A detectable marker is conjugated to a $Fc_\epsilon R$ molecule or a reagent in such a manner as not to block the ability of the $Fc_\epsilon R$ or reagent to bind to the IgE being
5 detected. Preferably, a carbohydrate group of a $Fc_\epsilon R$ is conjugated to biotin.

In another embodiment, a $Fc_\epsilon R$ molecule:IgE complex is detected by contacting a putative IgE-containing composition with a $Fc_\epsilon R$ molecule and then contacting the complex with an indicator molecule. Suitable indicator molecules of the present invention include molecules that can bind to either the $Fc_\epsilon R$ molecule or to the IgE
10 antibody. As such, an indicator molecule can comprise, for example, a $Fc_\epsilon R$ molecule, an antigen, an antibody and a lectin, depending upon which portion of the $Fc_\epsilon R$ molecule:IgE complex being detected. Preferred identifying labeled compounds that are antibodies include, for example, anti-IgE antibodies and anti- $Fc_\epsilon R$ antibodies. Preferred lectins include those lectins that bind to high-mannose groups. More preferred lectins
15 bind to high-mannose groups present on a $Fc_\epsilon R$ molecule of the present invention produced in insect cells. An indicator molecule itself can be attached to a detectable marker of the present invention. For example, an antibody can be conjugated to biotin, horseradish peroxidase, alkaline phosphatase or fluorescein.

In one preferred embodiment, a $Fc_\epsilon R$ molecule:IgE complex is detected by
20 contacting the complex with a reagent that selectively binds to a $Fc_\epsilon R$ molecule of the present invention. Examples of such a reagent includes, but are not limited to, an antibody that selectively binds to a $Fc_\epsilon R$ molecule (referred to herein as an anti- $Fc_\epsilon R$ antibody) or a compound that selectively binds to a detectable marker conjugated to a $Fc_\epsilon R$ molecule. $Fc_\epsilon R$ molecules conjugated to biotin are preferably detected using
25 streptavidin, more preferably using ImmunoPure® NeutrAvidin (available from Pierce, Rockford, IL).

In another preferred embodiment, a $Fc_\epsilon R$ molecule:IgE complex is detected by contacting the complex with a reagent that selectively binds to an IgE antibody (referred to herein as an anti-IgE reagent). Examples of such an anti-IgE reagent include, but are
30 not limited to, a secondary antibody that is an anti-isotype antibody (e.g., an antibody that selectively binds to the constant region of an IgE), an antibody-binding bacterial

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surface protein (e.g., Protein A or Protein G), an antibody-binding cell (e.g., a B cell, a T cell, a natural killer cell, a polymorphonuclear leukocyte cell, a monocyte cell or a macrophage cell), an antibody-binding eukaryotic cell surface protein (e.g., an Fc receptor), and an antibody-binding complement protein. Preferred anti-IgE reagents include, but are not limited to, D9, and CMI antibody #9, CMI antibody #19, CMI antibody #59 and CMI antibody #71 (available from Custom Monoclonal International, West Sacramento, CA). In particular, as used herein, an anti-IgE antibody includes not only a complete antibody but also any subunit or portion thereof that is capable of selectively binding to an IgE heavy chain constant region. For example, a portion of an anti-IgE reagent can include an Fab fragment or a F(ab')₂ fragment, which are described in detail in Janeway et al., in *Immunobiology, the Immune System in Health and Disease*, Garland Publishing, Inc., NY, 1996 (which is incorporated herein by this reference in its entirety).

In one embodiment a complex can be formed and detected in solution. In another embodiment, a complex can be formed in which one or more members of the complex are immobilized on (e.g., coated onto) a substrate. Immobilization techniques are known to those skilled in the art. Suitable substrate materials include, but are not limited to, plastic, glass, gel, celluloid, paper, PVDF (poly-vinylidene-fluoride), nylon, nitrocellulose, and particulate materials such as latex, polystyrene, nylon, nitrocellulose, agarose and magnetic resin. Suitable shapes for substrate material include, but are not limited to, a well (e.g., microtiter dish well), a plate, a dipstick, a bead, a lateral flow apparatus, a membrane, a filter, a tube, a dish, a celluloid-type matrix, a magnetic particle, and other particulates. A particularly preferred substrate comprises an ELISA plate, a dipstick, a radioimmunoassay plate, agarose beads, plastic beads, latex beads, immunoblot membranes and immunoblot papers. In one embodiment, a substrate, such as a particulate, can include a detectable marker.

A preferred method to detect IgE is an immunosorbent assay. An immunoabsorbent assay of the present invention comprises a capture molecule and an indicator molecule. A capture molecule of the present invention binds to an IgE in such a manner that the IgE is immobilized to a substrate. As such, a capture molecule is preferably immobilized to a substrate of the present invention prior to exposure of the

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capture molecule to a putative IgE-containing composition. An indicator molecule of the present invention detects the presence of an IgE bound to a capture molecule. As such, an indicator molecule preferably is not immobilized to the same substrate as a capture molecule prior to exposure of the capture molecule to a putative IgE-containing composition.

5 A preferred immunoabsorbent assay method includes a step of either: (a) binding an Fc_εR molecule to a substrate prior to contacting a Fc_εR molecule with a putative IgE-containing composition to form a Fc_εR molecule-coated substrate; or (b) binding a putative IgE-containing composition to a substrate prior to contacting a Fc_εR molecule
10 with a putative IgE-containing composition to form a putative IgE-containing composition-coated substrate. Preferably, the substrate includes of a non-coated substrate, a Fc_εR molecule-coated substrate, an antigen-coated substrate or an anti-IgE antibody-coated substrate.

Both a capture molecule and an indicator molecule of the present invention are
15 capable of binding to an IgE. Preferably, a capture molecule binds to a different region of an IgE than an indicator molecule, thereby allowing a capture molecule to be bound to an IgE at the same time as an indicator molecule. The use of a reagent as a capture molecule or an indicator molecule depends upon whether the molecule is immobilized to a substrate when the molecule is exposed to an IgE. For example, a Fc_εR molecule of
20 the present invention is used as a capture molecule when the Fc_εR molecule is bound to a substrate. Alternatively, a Fc_εR molecule is used as an indicator molecule when the Fc_εR molecule is not bound to a substrate. Suitable molecule for use as capture molecules or indicator molecules include, but are not limited to, a Fc_εR molecule of the present invention, an antigen reagent or an anti-IgE antibody reagent of the present
25 invention.

An immunoabsorbent assay of the present invention can further comprise one or more layers and/or types of secondary molecules or other binding molecules capable of detecting the presence of an indicator molecule. For example, an untagged (i.e., not conjugated to a detectable marker) secondary antibody that selectively binds to an
30 indicator molecule can be bound to a tagged (i.e., conjugated to a detectable marker) tertiary antibody that selectively binds to the secondary antibody. Suitable secondary

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antibodies, tertiary antibodies and other secondary or tertiary molecules can be selected by those of skill in the art. Preferred secondary molecules of the present invention include, an antigen, an anti-IgE idiotypic antibody and an anti-IgE isotypic. Preferred tertiary molecules can be selected by a skilled artisan based upon the characteristics of the secondary molecule. The same strategy is applied for subsequent layers.

In one embodiment, a desired antigen is used as a capture molecule by being immobilized on a substrate, such as a microtiter dish well or a dipstick. Preferred antigens include those disclosed herein. A biological sample collected from an animal is applied to the substrate and incubated under conditions suitable (i.e., sufficient) to allow for antigen:IgE complex formation bound to the substrate (i.e., IgE in a sample binds to an antigen immobilized on a substrate). Excess non-bound material (i.e., material from the biological sample that has not bound to the antigen), if any, is removed from the substrate under conditions that retain antigen:IgE complex binding to the substrate. Preferred conditions are disclosed herein in the Examples section and generally in Sambrook et al., *ibid*. An indicator molecule that can selectively bind to an IgE bound to the antigen, the indicator molecule can be conjugated to a detectable marker (preferably to an enzyme label, to a colorimetric label, to a fluorescent label, to a radioisotope, or to a ligand such as of the biotin or avidin family), is added to the substrate and incubated to allow formation of a complex between the indicator molecule and the antigen:IgE complex. Excess indicator molecule is removed, a developing agent is added if required, and the substrate is submitted to a detection device for analysis. A preferred indicator molecule for this embodiment is a Fc_εR molecule, preferably conjugated to biotin, to a fluorescent label or to an enzyme label.

In one embodiment, a Fc_εR molecule is used as a capture molecule by being immobilized on a substrate, such as a microtiter dish well or a dipstick. A biological sample collected from an animal is applied to the substrate and incubated under conditions suitable to allow for Fc_εR molecule:IgE complex formation bound to the substrate. Excess non-bound material, if any, is removed from the substrate under conditions that retain Fc_εR molecule:IgE complex binding to the substrate. An indicator molecule that can selectively bind to an IgE bound to the Fc_εR is added to the substrate and incubated to allow formation of a complex between the indicator molecule and the

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Fc_εR molecule:IgE complex. Preferably, the indicator molecule is conjugated to a detectable marker (preferably to an enzyme label, to a colorimetric label, to a fluorescent label, to a radioisotope, or to a ligand such as of the biotin or avidin family). Excess indicator molecule is removed, a developing agent is added if required, and the substrate is submitted to a detection device for analysis. A preferred indicator molecule for this embodiment is an antigen that will bind to IgE in the biological sample or an anti-IgE isotype or idiotype antibody, either preferably being conjugated to fluorescein or biotin.

In one embodiment, an anti-IgE antibody (e.g., isotype or idiotype specific antibody) is used as a capture molecule by being immobilized on a substrate, such as a microtiter dish well or a dipstick. A biological sample collected from an animal is applied to the substrate and incubated under conditions suitable to allow for anti-IgE antibody:IgE complex formation bound to the substrate. Excess non-bound material, if any, is removed from the substrate under conditions that retain anti-IgE antibody:IgE complex binding to the substrate. A Fc_εR molecule is added to the substrate and incubated to allow formation of a complex between the Fc_εR molecule and the anti-IgE antibody:IgE complex. Preferably, the Fc_εR molecule is conjugated to a detectable marker (preferably to biotin, an enzyme label or a fluorescent label). Excess Fc_εR molecule is removed, a developing agent is added if required, and the substrate is submitted to a detection device for analysis.

In one embodiment, an immunosorbent assay of the present invention does not utilize a capture molecule. In this embodiment, a biological sample collected from an animal is applied to a substrate, such as a microtiter dish well or a dipstick, and incubated under conditions suitable to allow for IgE binding to the substrate. Any IgE present in the bodily fluid is immobilized on the substrate. Excess non-bound material, if any, is removed from the substrate under conditions that retain IgE binding to the substrate. A Fc_εR molecule is added to the substrate and incubated to allow formation of a complex between the Fc_εR molecule and the IgE. Preferably, the Fc_εR molecule is conjugated to a detectable marker (preferably to biotin, an enzyme label or a fluorescent label). Excess Fc_εR molecule is removed, a developing agent is added if required, and the substrate is submitted to a detection device for analysis.

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Another preferred method to detect IgE is a lateral flow assay, examples of which are disclosed in U.S. Patent No. 5,424,193, issued June 13, 1995, by Pronovost et al.; U.S. Patent No. 5,415,994, issued May 16, 1995, by Imrich et al; WO 94/29696, published December 22, 1994, by Miller et al.; and WO 94/01775, published January 20, 1994, by Pawlak et al.; each of these patent publications is incorporated by reference herein in its entirety. In one embodiment, a biological sample is placed in a lateral flow apparatus that includes the following components: (a) a support structure defining a flow path; (b) a labeling reagent comprising a bead conjugated to an antigen, the labeling reagent being impregnated within the support structure in a labeling zone; and (c) a capture reagent comprising an IgE-binding composition. Preferred antigens include those disclosed herein. The capture reagent is located downstream of the labeling reagent within a capture zone fluidly connected to the labeling zone in such a manner that the labeling reagent can flow from the labeling zone into the capture zone. The support structure comprises a material that does not impede the flow of the beads from the labeling zone to the capture zone. Suitable materials for use as a support structure include ionic (i.e., anionic or cationic) material. Examples of such a material include, but are not limited to, nitrocellulose (NC), PVDF, carboxymethylcellulose (CM). The support structure defines a flow path that is lateral and is divided into zones, namely a labeling zone and a capture zone. The apparatus can further comprise a sample receiving zone located along the flow path, more preferably upstream of the labeling reagent. The flow path in the support structure is created by contacting a portion of the support structure downstream of the capture zone, preferably at the end of the flow path, to an absorbent capable of absorbing excess liquid from the labeling and capture zones.

In this embodiment, the biological sample is applied to the sample receiving zone which includes a portion of the support structure. The labeling zone receives the sample from the sample receiving zone which is directed downstream by the flow path. The labeling zone comprises the labeling reagent that binds to IgE. A preferred labeling reagent is an antigen conjugated, either directly or through a linker, to a plastic bead substrate, such as to a latex bead. The substrate also includes a detectable marker, preferably a colorimetric marker. Typically, the labeling reagent is impregnated to the support structure by drying or lyophilization. The sample structure also comprises a

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capture zone downstream of the labeling zone. The capture zone receives labeling reagent from the labeling zone which is directed downstream by the flow path. The capture zone contains the capture reagent, in this case a $Fc_\epsilon R$ molecule, as disclosed above, that immobilizes the IgE complexed to the antigen in the capture zone. The capture reagent is preferably fixed to the support structure by drying or lyophilizing. The labeling reagent accumulates in the capture zone and the accumulation is assessed visually or by an optical detection device.

In another embodiment, a lateral flow apparatus used to detect IgE includes: (a) a support structure defining a flow path; (b) a labeling reagent comprising a $Fc_\epsilon R$ molecule as described above, the labeling reagent impregnated within the support structure in a labeling zone; and (c) a capture reagent comprising an antigen, the capture reagent being located downstream of the labeling reagent within a capture zone fluidly connected to the labeling zone in such a manner that the labeling reagent can flow from the labeling zone into the capture zone. The apparatus preferably also includes a sample receiving zone located along the flow path, preferably upstream of the labeling reagent. The apparatus preferably also includes an absorbent located at the end of the flow path.

One embodiment of the present invention is an inhibition assay in which the presence of IgE in a putative IgE-containing composition is determined by adding such composition to a $Fc_\epsilon R$ molecule of the present invention and an isolated IgE known to bind to the $Fc_\epsilon R$ molecule. The absence of binding of the $Fc_\epsilon R$ molecule to the known IgE indicating the presence of IgE in the putative IgE-containing composition.

The present invention also includes kits to detect IgE based on each of the disclosed detection methods. One embodiment is a kit to detect IgE comprising a human Fc_ϵ receptor ($Fc_\epsilon R$) molecule and a means for detecting an IgE including canine IgE, feline IgE and/or equine IgE. Suitable and preferred $Fc_\epsilon R$ molecules are disclosed herein. Suitable means of detection include compounds disclosed herein that bind to either the $Fc_\epsilon R$ molecule or to an IgE. A preferred kit of the present invention further comprises a detection means including one or more antigens disclosed herein, an antibody capable of selectively binding to an IgE disclosed herein and/or a compound capable of binding to a detectable marker conjugated to a $Fc_\epsilon R$ molecule (e.g., avidin, streptavidin and ImmunoPure® NeutrAvidin when the detectable marker is biotin).

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Such antigens preferably induce IgE antibody production in animals including canines, felines and/or equines.

A preferred embodiment of a kit of the present invention is a flea allergen kit comprising a flea allergen such as those disclosed herein. A particularly preferred flea
5 allergen for use with a flea allergen kit includes a flea saliva product or a flea saliva protein.

Another preferred kit of the present invention is a general allergen kit comprising an allergen common to all regions of the United States and a human $Fc\epsilon R$ molecule of the present invention. As used herein, a "general allergen" kit refers to a kit comprising
10 allergens that are found substantially throughout the United States (i.e., essentially not limited to certain regions of the United States). A general allergen kit provides an advantage over regional allergen kits because a single kit can be used to test an animal located in most geographical locations on the United States. Suitable and preferred general allergens for use with a general allergen kit of the present invention include
15 those general allergens disclosed herein.

Another preferred kit of the present invention is a food allergen kit comprising a food allergen including beef, chicken, pork, a mixture of fish, such as cod, halibut or and tuna, egg, milk, Brewer's yeast, whole wheat, corn, soybean, cheese and rice, and a
20 human $Fc\epsilon R$ molecule of the present invention. Preferably, the beef, chicken, pork, fish, corn and rice, are cooked.

A preferred kit of the present invention includes those in which the allergen is immobilized to a substrate. If a kit comprises two or more antigens, the kit can comprise one or more compositions, each composition comprising one antigen. As such, each antigen can be tested separately. A kit can also contain two or more diagnostic reagents
25 for IgE, additional isolated IgE antigens and/or antibodies as disclosed herein.

Particularly preferred are kits used in a lateral flow assay format. It is within the scope of the present invention that a lateral flow assay kit can include one or more lateral flow assay apparatuses. Multiple lateral flow apparatuses can be attached to each other at one end of each apparatus, thereby creating a fan-like structure.

30 In particular, a method and kit of the present invention are useful for diagnosing abnormal conditions in animals that are associated with changing levels of IgE.

Particularly preferred conditions to diagnose include allergies, parasitic infections and neoplasia. For example, a method and kit of the present invention are particularly useful for detecting flea allergy dermatitis (FAD), when such method or kit includes the use of a flea saliva antigen. FAD is defined as a hypersensitive response to fleabites.

- 5 Preferably, a putative IgE-containing composition is obtained from an animal suspected of having FAD. Preferred animals include those disclosed herein, with dogs and cats being more preferred. In addition, methods and kits of the present invention are particularly useful for detecting helminth infection, in particular heartworm infection, when such methods or kits include the use of a helminth antigen, such as Di33.
- 10 Preferably, a putative IgE-containing composition is obtained from an animal suspected of having a helminth infection. Preferred animals include those disclosed herein, with dogs and cats being more preferred.

The following examples are provided for the purposes of illustration and are not intended to limit the scope of the present invention.

15 Examples

Example 1.

This example describes the construction of a recombinant baculovirus expressing a truncated portion of the α -chain of the human Fc_ϵ receptor.

- Recombinant molecule pVL-nhFc $_\epsilon$ R α_{612} , containing a nucleic acid molecule
- 20 encoding the extracellular domain of the Fc $_\epsilon$ R α chain, operatively linked to baculovirus polyhedron transcription control sequences was produced in the following manner. A cDNA clone encoding the full-length alpha chain (α chain) of the human Fc $_\epsilon$ receptor was obtained from Dr. Jean-Pierre Kinet (Harvard University, Cambridge, MA). The cDNA clone included an about 1198 nucleotide insert, referred to herein as nhFc $_\epsilon$ R α_{1198} .
- 25 The nucleic acid sequence of the coding strand of nhFc $_\epsilon$ R α_{1198} is denoted herein as SEQ ID NO:1. Translation of SEQ ID NO:1 indicates that nucleic acid molecule nhFc $_\epsilon$ R α_{1198} encodes a full-length human Fc $_\epsilon$ receptor α chain protein of about 257 amino acids, referred to herein as PhFc $_\epsilon$ R α_{257} , having amino acid sequence SEQ ID NO:2, assuming an open reading frame in which the initiation codon spans from nucleotide 107 through
- 30 nucleotide 109 of SEQ ID NO:1 and the termination codon spans from nucleotide 878 through nucleotide 880 of SEQ ID NO:1. The complement of SEQ ID NO:1 is

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represented herein by SEQ ID NO:3. The proposed mature protein (i.e., Fc_εRα chain from which the signal sequence has been cleaved), denoted herein as PhFc_εRα₂₃₂, contains about 232 amino acids which is represented herein as SEQ ID NO:6. The nucleic acid molecule encoding PhFc_εRα₂₃₂ is denoted herein as nhFc_εRα₆₉₉, the coding strand of which is represented by SEQ ID NO:7.

To produce a secreted form of the extracellular domain of the Fc_εR α chain, the hydrophobic transmembrane domain and the cytoplasmic tail of the Fc_εR α chain encoded by nhFc_εRα₁₁₉₈ were removed as follows. A Fc_εR α chain extracellular domain nucleic acid molecule-containing fragment of about 612 nucleotides was PCR amplified from nhFc_εRα₁₁₉₈ using a forward primer EJH 040 containing a *Bam*HI site, having the nucleic acid sequence 5' CGC GGA TCC TAT AAA TAT GGC TCC TGC CAT GG 3' (denoted SEQ ID NO:8) and a reverse primer IgE ANTI-SENSE containing an *Eco*RI site, having the nucleic acid sequence 5' GGC GAA TTC TTA AGC TTT TAT TAC AG 3' (denoted herein as SEQ ID NO:9). The resulting PCR product was digested with *Bam*HI and *Eco*RI to produce nhFc_εRα₆₁₂. Nucleic acid molecule nhFc_εRα₆₁₂ contained an about 591 nucleotide fragment encoding the extracellular domain of the human Fc_εR α chain, extending from nucleotide 107 to nucleotide 697 of SEQ ID NO 1, denoted herein as nucleic acid molecule nhFc_εRα₅₉₁, the coding strand of which has a nucleic acid sequence denoted SEQ ID NO:10. Translation of SEQ ID NO:10 indicates that nucleic acid molecule nhFc_εRα₆₁₂ encodes a Fc_εR protein of about 197 amino acids, referred to herein as PhFc_εRα₁₉₇, having amino acid sequence SEQ ID NO:11. Nucleic acid molecule nhFc_εRα₆₁₂ encodes a secretable form of the human Fc_εR α chain which does not possess a leader sequence, which is denoted herein as PhFc_εRα₁₇₂ having amino acid sequence SEQ ID NO:13. The coding region for PhFc_εRα₁₇₂ is denoted nhFc_εRα₅₁₆, the coding strand of which has a nucleic acid sequence denoted SEQ ID NO:12. The complement of SEQ ID NO:12 is represented herein by SEQ ID NO:14.

In order to produce a baculovirus recombinant molecule capable of directing the production of PhFc_εRα₁₉₇, the nucleic acid molecule nhFc_εRα₆₁₂ was subcloned into unique *Bam*HI and *Eco*RI sites of pVL1392 baculovirus shuttle plasmid (available from Pharmingen, San Diego, CA) to produce a recombinant molecule referred to herein as

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pVL-nhFc_εRα₆₁₂. The resultant recombinant molecule pVL-nhFc_εRα₆₁₂ was verified for proper insert orientation by restriction mapping.

Example 2.

This example describes the production of PhFc_εRα₁₇₂ protein.

- 5 The recombinant molecule pVL-nhFc_εRα₆₁₂ was co-transfected with a linear Baculogold baculovirus DNA (available from Pharmingen) into *Trichoplusia ni* cells (available from Invitrogen Corp., San Diego, CA; High Five™ cells) using the following method. About 1.5 liter cultures of serum-free ex-Cell Medium (available from Invitrogen) were seeded with about 1 x 10⁶ cells per ml of medium. The *Trichoplusia ni*
- 10 cells were infected with recombinant molecule pVL-nhFc_εRα₆₁₂ at a multiplicity of infection (MOI) of about 2 to about 5 particle forming units (pfu) per cell to produce recombinant cell *Trichoplusia ni*-pVL-nhFc_εRα₆₁₂. The infection was allowed to proceed at a controlled temperature of 27°C for 48 hours, to produce recombinant protein PhFc_εRα₁₇₂. Following infection, cells were separated from the medium by
- 15 centrifugation, and the medium was frozen at -70°C.

- PhFc_εRα₁₇₂ was purified from the culture medium described immediately above by affinity chromatography using an IgE antibody produced by the myeloma cell line U266DI (American Tissue Type Catalogue No. TIB196) linked to sepharose 4B. The amino acid composition and N-terminal amino acid sequence of the affinity purified
- 20 PhFc_εRα₁₇₂ were determined using methods standard in the art. The results indicated that PhFc_εRα₁₇₂ was properly processed by the *Trichoplusia ni* cells.

Example 3.

This example describes the biotinylation of a recombinant human Fc_εR alpha chain protein.

- 25 Affinity purified recombinant protein PhFc_εRα₁₇₂, prepared as described above in Example 2, was biotinylated as follows. About 440 micrograms (μg) of PhFc_εRα₁₇₂ were diluted in about 1.5 milliliter (ml) of acetate buffer (0.1 M NaAc, pH 5.5) containing about 200 microliter (μl) of 0.1 M NaIO₄. The mixture was incubated for about 20 minutes, on ice, and about 2 μl of glycerol was added following the incubation.
- 30 The mixture was then dialyzed against about 2 liters of acetate buffer in a 3 ml Slide-A-Lyzer cassette (available from Pierce, Rockford, IL), 2 times for about 2 hours each

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time. About 3.72 μ g of biotin-LC-hydrazide (available from Pierce) was dissolved in about 200 μ l of dimethylsulfoxide (DMSO) and injected into the cassette. The cassette was then rocked at room temperature for about 2 hours. Following the incubation, the mixture containing recombinant protein and biotin was dialyzed 3 times, a first time for
5 about 18 hours and two times for about 2 hours, each time at 5°C against phosphate buffered saline. The biotinylated protein was recovered from the dialysis, and is referred to herein as PhFc ϵ R α ₁₇₂-BIOT.

Example 4.

This example describes detection of canine IgE in a solid-phase ELISA using
10 PhFc ϵ R α ₁₇₂-BIOT.

Wells of two Immulon II microtiter plates (available from Dynatech, Alexandria, VA) were coated with duplicate samples of about 100 μ l/well of various concentrations of purified canine IgE as denoted in Fig. 1. The canine IgE was obtained from a canine IgE producing hybridoma, such as heterohybridoma 2.39 (described in Gebhard et al.,
15 *Immunology* 85:429-434, 1995) and was diluted in a CBC buffer (15 mM Na₂CO₃ and 34.8 mM NaHCO₃, pH 9.6. The coated plates were incubated overnight at 4°C.

Following incubation, the canine IgE-containing solution was removed from each plate, and the plates were blotted dry. The plates were then blocked using about 200 μ l/well of 0.25% bovine serum albumin (BSA) contained in phosphate buffered saline (PBSB) for
20 about 1 hour at room temperature. The plates were then washed four times with 0.05% Tween-20 in PBS (PBST) using an automatic washer (available from Dynatech).

Experimental samples consisting of about 100 μ l/well of a 1:4000 dilution of 40 μ g/ml PhFc ϵ R α ₁₇₂-BIOT (about 145 μ g/ml; described in Example 3), contained in PBSB with 0.05% Tween-20 (PBSBT) were added to each well of one plate coated with canine IgE.

25 Control samples consisting of about 100 μ l of biotinylated anti-canine IgE monoclonal antibody D9 (supplied by Dr. DeBoer, U. of Wisconsin, Madison, WI) were added to each well of the other plate coated with canine IgE. The plates were incubated for 1 hour at room temperature and then washed four times with PBST. About 100 μ l of about 0.25 μ g/ml streptavidin conjugated to horseradish peroxidase (available from
30 Kirkegaard and Perry Laboratories (KPL), Gaithersburg, MD; diluted in PBST) was added to each well that received experimental or control samples. The plates were then

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incubated for 1 hour at room temperature and washed four times with PBST. About 100 μ l of TMB substrate (available from available from KPL), that had been pre-warmed to room temperature, was added. Plates were then incubated for 10 minutes at room temperature and then about 100 μ l/well of Stop Solution (available from KPL) was
5 added. Optical densities of wells were read on a Spectramax Microtiter Plate (available from Molecular Devices Inc.) reader at 450 nm within 10 minutes of adding the stop solution.

The results shown in Fig. 1 indicate that the alpha chain of human $\text{Fc}_\epsilon\text{R}$ detects the presence of canine IgE (closed circles) in a solid-phase assay in a similar manner as
10 the control antibody that binds specifically to canine IgE (D9; open circles).

Example 5.

This example describes detection of plant allergen-specific canine IgE using $\text{PhFc}_\epsilon\text{R}\alpha_{172}$ -BIOT.

Multiple wells of an Immulon II microtiter plate (available from Dynatech) were
15 coated with either about 100 μ l/well of 1 μ g/ml of Kentucky Blue Grass allergen or about 100 μ l/well of about 1 μ g/ml of Green Ash allergen (both available from Greer Inc., Lenoir, NC) both diluted in CBC buffer. The plate was incubated overnight at 4°C. The plate was blocked and washed as described in Example 4. Two different pools of canine sera were then added to the antigen-coated wells. The first pool consisted of sera
20 isolated from 8 dogs reported to be allergen reactive. The second pool consisted of sera isolated from 8 dogs reported to be allergen non-reactive. Each pool of sera was diluted 1:10 or 1:100 in PBST. About 100 μ l of each concentration of each diluted sera sample was added to the wells and incubated for 1 hour at room temperature. The plate was then washed four times with PBST. About 100 μ l/well of a 1:4000 dilution of 40 μ g/ml
25 $\text{PhFc}_\epsilon\text{R}\alpha_{172}$ -BIOT (described in Example 3), contained in PBSBT was added to the antigen-coated wells. The plate was incubated for 1 hour at room temperature. The plate was then washed four times with PBST. About 100 μ l/well of about 0.25 μ g/ml of neutravidin conjugated to horseradish peroxidase (available from Pierce) contained in PBSBT, was added. The plate was incubated for 1 hour at room temperature. The plate
30 was then washed and the presence of neutravidin bound to the plate detected using the method described in Example 4.

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The results shown in Fig. 2 indicate that the alpha chain of human $Fc_\epsilon R$ detects the presence of canine IgE antibodies that bind specifically to a common grass allergen or to a common tree allergen. In addition, detection of canine IgE antibodies is dose dependent.

5 Example 6.

This example describes detection of total canine IgE using $PhFc_\epsilon R\alpha_{172}$ -BIOT.

Multiple wells of an Immulon II microtiter plate (available from Dynatech) were coated with about 100 μ l/well of about 1 μ g/ml CMI anti-canine IgE antibody #6 (available from Custom Monoclonals International, West Sacramento, CA) diluted in
10 CBC buffer. The plate was incubated overnight at 4°C. The plate was blocked and washed as described in Example 4. About 100 μ l/well of a 1:60 dilution in PBSBT of sera samples from a variety of sources were then added to multiple wells coated with anti-IgE antibody. The samples included: (1) serum from a dog known to be allergic to flea saliva; (2) serum from dogs infected with *D. immitis*; (3) and (4) a pool of dog sera
15 from defined as canine allergy calibrators (available from BioProducts DVM, Tempe, AZ); (5) pools of dog sera containing antibodies that have low binding to Kentucky Blue Grass allergen; (6) pools of dog sera that have high binding to Kentucky Blue Grass allergen; (7) a pool of dog sera from dogs known to be allergic to flea saliva, the sample was heat inactivated (at 56°C for 4 hours); (8) a pool of dog sera from dogs
20 known to be allergic to flea saliva; or (9) a pool of dog sera from dogs raised in a barrier facility (i.e., negative control). A set of positive control samples consisting of IgE derived from the canine heterohybridoma described in Example 4 were also added to the plate to generate a standard curve. The plate was incubated for 1 hour at room temperature and then washed four times with PBST. The presence of canine IgE was
25 detected using either about 100 μ l/well of a 1:4000 dilution of 40 μ g/ml $PhFc_\epsilon R\alpha_{172}$ -BIOT (described in Example 3) or about 100 μ l/well of about 1 μ g/ml CMI anti-canine IgE antibody #19 (available from Custom Monoclonals International), both contained in PBSBT. The plate was incubated for 1 hour at room temperature. The plate was then washed, contacted with about 0.25 μ g/ml streptavidin conjugated to horseradish
30 peroxidase, washed again, and the presence of streptavidin bound to the plate was detected using the method described in Example 4. The optical density readings

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obtained for the control samples were used to generate a standard curve that was used to determine the total IgE bound to wells that had received test samples.

The results shown in Fig. 3 indicate that canine IgE from a variety of dog sera are detected using the alpha chain of human $Fc_\epsilon R$ in a manner similar to using an antibody that binds specifically to canine IgE. The absence of detectable amounts of IgE in the heat treated sample (Sample 7) indicates that the antibody detected by $PhFc_\epsilon R\alpha_{172}$ -BIOT is IgE. In addition, the results indicate that $PhFc_\epsilon R\alpha_{172}$ -BIOT is an effective reagent for detecting IgE that binds to allergen Kentucky Blue Grass, Samples 5 and 6), as well as a parasite antigen (*D. Immitis*, Sample 2).

10 Example 7.

This example describes detection of canine IgE in dog sera isolated from dogs known to be allergic to flea saliva, using $PhFc_\epsilon R\alpha_{172}$ -BIOT.

Multiple wells of an Immulon II microtiter plate were coated with about 100 μ l/well of varying concentrations of flea saliva recombinant protein fspN (described in PCT Patent Publication No. WO 96/11271, *ibid.*; concentrations shown in Fig. 4) diluted in CBC buffer. The plate was incubated overnight at 4°C. The plate was then blocked and washed as described in Example 4. About 100 μ l/well of a 1:10 dilution in PBSBT of a pool of sera isolated from dogs known to produce IgE that binds specifically to flea saliva. Some wells did not receive dog sera so that background binding levels could be determined. The plate was incubated for 1 hour at room temperature and then washed four times with PBST. About 100 μ l/well of a 1:4000 dilution of 40 μ g/ml $PhFc_\epsilon R\alpha_{172}$ -BIOT (described in Example 3) contained in PBSBT was added. The plate was incubated for 1 hour at room temperature. The plate was then washed, contacted with about 0.25 μ g/ml streptavidin-conjugated to horseradish peroxidase, washed again, and the presence of streptavidin bound to the plate was detected using the method described in Example 4.

The results shown in Fig. 4 indicate that canine IgE that binds specifically to a flea saliva antigen is detected using the alpha chain of human $Fc_\epsilon R$.

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Example 8.

This example describes detection of total canine IgE in dog sera isolated from dogs known to be allergic to flea saliva, heartworm-infected dogs and specific pathogen free (SPF) dogs, using PhFc_εRα₁₇₂-BIOT.

- 5 Multiple wells of an Immulon II microtiter plate were coated with about 100 μl/well of about 1 μg/ml CMI anti-canine IgE antibody #6 (available from Custom Monoclonals International) in CBC buffer. The plate was incubated overnight at 4°C. The plate was blocked and washed as described in Example 4. About 100 μl/well of different samples of IgE-containing fluids in PBSBT were added to multiple wells
- 10 coated with the anti-canine IgE antibody. The samples included: (1) 100 μg/ml of canine IgE purified from the heterohybridoma described in Example 4; (2) a 1:10 dilution of a pool of sera from dogs known to be allergic to flea saliva, (3) a 1:10 dilution of the same sera pool as in (2) but heat inactivated; (4) a 1:10 dilution of serum from a dog known to have clinical flea allergy dermatitis (dog CPO2); (5) a 1:10 dilution
- 15 of heat inactivated CPO2 serum; (6) a 1:10 dilution of serum from a heartworm-infected dog (dog 417); (7) a 1:10 dilution of heat inactivated 417 serum; (8) a 1:10 dilution of a pool of sera from heartworm-infected dogs; (9) a 1:10 dilution of the same sera pool as in (8) but heat inactivated; and (10) a pool of sera from dogs raised in a barrier facility. Each sample was diluted in PBSBT. The plate was incubated for 1 hour at room
- 20 temperature and then washed four times with PBST. About 100 μl/well of a 1:4000 dilution of 40 μg/ml PhFc_εRα₁₇₂-BIOT (described in Example 3) in PBSBT was added. The plate was incubated for 1 hour at room temperature. The plate was then washed, contacted with about 0.25 ug/ml streptavidin-conjugated to horseradish peroxidase, washed again, and the presence of streptavidin bound to the plate was detected using the
- 25 method described in Example 4.

- The results shown in Fig. 5 indicate that canine IgE from dogs allergic to flea saliva and from dogs infected with heartworm are detected using the alpha chain of human Fc_εR. In addition, the absence of colorimetric signal in samples of heat inactivated sera indicates that antibody bound to the anti-IgE antibody and detected by
- 30 Fc_εR alpha chain is an epsilon isotype antibody and not another isotype.

Example 9.

This example describes detection of IgE that specifically binds to flea saliva, using PhFc_εRα₁₇₂-BIOT.

Multiple wells of an Immulon II microtiter plate were coated with about 100 μl/well of about 0.1 μg/ml of flea saliva collected using the method described in PCT Patent Publication No. WO 96/11271, *ibid.*, in CBC buffer. The plate was incubated, blocked and washed as described in Example 4. The IgE-containing samples described in Example 8 were then applied to the flea saliva coated plate. The plate was then treated using the method described in Example 8.

The results shown in Fig. 6 indicate that canine IgE that binds specifically to flea saliva, contained in serum, is detected using the alpha chain of human Fc_εR. In addition, the absence of colorimetric signal in samples of heat inactivated serum indicates that antibody bound to the flea saliva protein and detected by Fc_εR alpha chain is an epsilon isotype antibody.

Example 10.

This example describes the detection of feline IgE using PhFc_εRα₁₇₂-BIOT.

Multiple wells of an Immulon II microtiter plate were coated with about 100 μl/well of about 10 μg/ml Di33 protein (described in U.S. Patent Application Serial No. 08/715,628, *ibid.*) or 10 μg/ml crude homogenate of heartworm, both in CBC buffer. Crude homogenate of heartworm is the clarified supernatant of adult heartworms homogenized in PBS. The plate was incubated overnight at 4°C. The plate was blocked and washed as described in Example 4. Serum samples from 2 heartworm infected cats were then added to Di33-coated wells and to heartworm antigen-coated wells. About 100 μl/well of a 1:10 dilution in PBSBT of sera from heartworm-infected cat # AXH3 or from cat #MGC2 were added to the plate. Negative control samples consisting of serum from pre-infection bleeds of cat #AXH3 and cat# MGC2 were also added to the plate at a dilution of 1:10 in PBSBT. A positive control sample consisting of a pool of sera from heartworm-infected dogs was also added to the plate at a dilution of 1:10 in PBSBT. The plate was incubated for 1 hour at room temperature and then washed four times with PBST. About 100 μl/well of a 1:4000 dilution of 40 μg/ml PhFc_εRα₁₇₂-BIOT (described in Example 3) in PBSBT was added. The plate was incubated for 1 hour at room

temperature. The plate was then washed, contacted with 1:4000 dilution of a 0.5 mg/ml solution of streptavidin-conjugated to horseradish peroxidase, washed again, and the presence of streptavidin bound to the plate was detected using the method described in Example 4.

- 5 The results shown in Fig. 7 indicate that feline IgE that binds specifically to crude homogenate of heartworm or Di33 protein is detected using the alpha chain of human $Fc_\epsilon R$.

Example 11.

This example describes detection of feline IgE using $PhFc_\epsilon R\alpha_{172}$ -BIOT.

- 10 Multiple wells of an Immulon II microtiter plate were coated with Di33 as described in Example 10, in CBC buffer. The plate was incubated overnight at 4°C. The plate was blocked and washed as described in Example 4. Serum samples from 2 heartworm infected cats were then added to Di33-coated wells. About 100 µl/well of a 1:10 dilution in PBSBT of serum from heartworm-infected cat # MGC2 and a pool of
15 sera from heartworm-infected cats, as well as heat inactivated samples of each of these sera, were added to the plate. A positive control sample consisting of a pool of sera from heartworm-infected dogs was also added to the plate at a dilution of 1:10 in PBSBT. The plate was incubated for 1 hour at room temperature and then washed four times with PBST. About 100 µl/well of a 1:4000 dilution of 40 µg/ml $PhFc_\epsilon R\alpha_{172}$ -BIOT
20 (described in Example 3) in PBSBT was added. The plate was incubated for 1 hour at room temperature. The plate was then washed, contacted with streptavidin-conjugated to horseradish peroxidase, washed again, and the presence of streptavidin bound to the plate was detected using the method described in Example 4.

- 25 The results shown in Fig. 8 indicate that feline IgE from heartworm-infected cats that specifically binds to the heartworm antigen Di33 is detected using the alpha chain of human $Fc_\epsilon R$. In addition, the absence of colorimetric signal in samples of heat inactivated sera indicates that antibody bound to the Di33 protein and detected by $Fc_\epsilon R$ alpha chain is an epsilon isotype antibody.

Example 12

- 30 This example describes detection of equine IgE in a solid-phase ELISA using $PhFc_\epsilon R\alpha_{172}$ -BIOT.

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Horse sera from a horse known to be allergic to certain allergens and horse sera from a horse known not to be allergic the same allergens, were assayed for the presence of IgE using PhFc_εRα₁₇₂-BIOT as follows. A North Atlantic/Ohio Valley Regional Panel plate of a Canitec™ Allergen-Specific IgE Kit (available from BioProducts DVM) was blocked and washed as described in Example 4. Two samples of about 1:10 dilutions of the two horse sera were prepared using PBSBT. The two samples were added to the blocked plate and the plate was incubated for 1 hour at room temperature. The plate was washed as described in Example 4. About 100 μl/well of a 1:4000 dilution of 40 μg/ml PhFc_εRα₁₇₂-BIOT (described in Example 3), contained in PBSBT was added to each well. The plate was then washed, contacted with 1:4000 dilution of a 0.5 mg/ml solution of streptavidin-conjugated to horseradish peroxidase, washed again, and the presence of streptavidin bound to the plate was detected using the method described in Example 4.

The results shown in Fig. 9 indicate that equine IgE from a horse known to be allergic to certain allergens specifically binds to certain plant and mite allergens is detected using the alpha chain of human Fc_εR.

Example 13

This example describes detection of canine IgE in a solid-phase ELISA using basophilic cells transfected with human Fc_εR alpha chain.

Rat basophilic leukemia (RBL) cells transfected with a nucleic acid molecule encoding a human Fc_εR alpha chain (referred to herein as RBL-hFc_εR cells; described in Miller et al., *Science* 244:334-337, 1989) were used to detect canine IgE as follows. About 4 x 10⁴ RBL-hFc_εR cells contained in Earles Modified Eagles Medium containing 10% fetal bovine serum (EMEM-FBS) were added to each well of 96-well flat bottom tissue culture plates. The RBL-hFc_εR cells were incubated overnight at 37°C. Following the incubation the plates were washed 4 times with PBST. The cells were then fixed for about 2 minutes using about 200 μl per well of absolute alcohol at room temperature. The plates were then washed 8 times with PBST to remove residual alcohol.

Serial dilutions in EMEM-FBS (concentrations shown in Fig. 10) were prepared using a pool of sera from dogs infected with heartworm. Serial dilutions in EMEM-FBS

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(concentrations shown in Fig. 11) were prepared using a pool of sera from dogs sensitized to flea saliva. Additional samples were prepared in which both pools of sera were heat inactivated for about 4-hours at 56°C. The heat treated samples were diluted as described above.

- 5 About 100 µl of each dilution of each serum sample was added to separate wells containing fixed RBL-hFc_εR cells and the plates were incubated at 37°C for about 1 hour. Following the incubation, the plates were washed 4 times with PBST. About 5 µg of a murine IgG monoclonal antibody anti-canine IgE antibody (i.e., Custom Monoclonal Antibody #71; available from Custom Monoclonal International) in 100 µl
10 of EMEM-FBS was added to each well. The plates were incubated for about 30 minutes at 37°C. Following the incubation, the plates were washed 4 times with PBST. About 100 ng of horseradish peroxidase labelled donkey anti-murine IgG (available from Jackson Laboratories, Westgrove, PA) in 100 µl of EMEM-FBS was added to each well, and the plates were incubated for about 30 minutes at room temperature. Following the
15 incubation, the plates were washed 4 times with PBST. The presence of anti-murine IgG bound to the plates thereby indicating the ability of RBL-hFc_εR cells to bind to canine IgE was detected using the method described in Example 4.

- The results shown in Fig. 10 indicate that canine IgE from heartworm-infected dogs (◆) is detected using RBL-h Fc_εR cells expressing the alpha chain of human Fc_εR.
20 In addition, the absence of colorimetric signal in samples of heat inactivated samples of such sera (■) indicates that antibody detected by the Fc_εR alpha chain on the RBL-h Fc_εR cells is an epsilon isotype antibody. Similarly, the results shown in Fig. 11 indicate that canine IgE from dogs sensitized with flea saliva (◆) is detected using RBL-h Fc_εR cells expressing the alpha chain of human Fc_εR. In addition, the absence of colorimetric
25 signal in samples of heat inactivated samples of such sera (■) indicates that antibody detected by the Fc_εR alpha chain on the RBL-h Fc_εR cells is an epsilon isotype antibody.

-35-

SEQUENCE LISTING

(1) GENERAL INFORMATION:

- (i) APPLICANT:
5 (A) NAME: Heska Corporation
(B) STREET: 1825 Sharp Point Drive
(C) CITY: Fort Collins
(D) STATE: CO
(E) COUNTRY: US
10 (F) POSTAL CODE (ZIP): 80525
(G) TELEPHONE: (970) 493-7272
(H) TELEFAX: (970) 484-9505
- (ii) TITLE OF INVENTION: METHOD TO DETECT IGE
- (iii) NUMBER OF SEQUENCES: 13
- (iv) CORRESPONDENCE ADDRESS:
15 (A) ADDRESSEE: LAHIVE & COCKFIELD, LLP
(B) STREET: 28 STATE STREET
(C) CITY: BOSTON
(D) STATE: MA
20 (E) COUNTRY: US
(F) ZIP: 02109
- (v) COMPUTER READABLE FORM:
25 (A) MEDIUM TYPE: Floppy disk
(B) COMPUTER: IBM PC compatible
(C) OPERATING SYSTEM: Windows 95
(D) SOFTWARE: ASCII DOS TEXT
- (vi) CURRENT APPLICATION DATA:
(A) APPLICATION NUMBER: 08/756,387
(B) FILING DATE: November 26, 1996
(C) CLASSIFICATION:
- 30 (vii) PRIOR APPLICATION DATA:
(A) APPLICATION NUMBER:
(B) FILING DATE:
- (viii) ATTORNEY/AGENT INFORMATION:
35 (A) NAME: Rothenberger, Scott D.
(B) REGISTRATION NUMBER: 41,277
(C) REFERENCE/DOCKET NUMBER:
- (ix) TELECOMMUNICATION INFORMATION:
(A) TELEPHONE: (617) 227-7400
(B) TELEFAX: (617) 742-4214

40 (2) INFORMATION FOR SEQ ID NO:1:

- (i) SEQUENCE CHARACTERISTICS:
45 (A) LENGTH: 1198 nucleotides
(B) TYPE: nucleic acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: cDNA
- (iii) FEATURE:
(A) NAME/KEY: CDS
(B) LOCATION: 107..877

-36-

(iv) SEQUENCE DESCRIPTION: SEQ ID NO:1

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	20 25 30 35						
	CCT CCA TGG AAT AGA ATA TTT AAA GGA GAG AAT GTG ACT CTT ACA TGT	259					
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	55 60 65						
	GGC AGC CTT TCA GAA GAG ACA AAT TCA AGT TTG AAT ATT GTG AAT GCC	355					
	Gly Ser Leu Ser Glu Glu Thr Asn Ser Ser Leu Asn Ile Val Asn Ala						
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	Lys Phe Glu Asp Ser Gly Glu Tyr Lys Cys Gln His Gln Gln Val Asn						
	85 90 95						
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	Glu Ser Glu Pro Val Tyr Leu Glu Val Phe Ser Asp Trp Leu Leu Leu						
	100 105 110 115						
	CAG GCC TCT GCT GAG GTG GTG ATG GAG GGC CAG CCC CTC TTC CTC AGG	499					
	Gln Ala Ser Ala Glu Val Val Met Glu Gly Gln Pro Leu Phe Leu Arg						
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	Cys His Gly Trp Arg Asn Trp Asp Val Tyr Lys Val Ile Tyr Tyr Lys						
	135 140 145						
	GAT GGT GAA GCT CTC AAG TAC TGG TAT GAG AAC CAC AAC ATC TCC ATT	595					
	Asp Gly Glu Ala Leu Lys Tyr Trp Tyr Glu Asn His Asn Ile Ser Ile						
35	150 155 160						
	ACA AAT GCC ACA GTT GAA GAC AGT GGA ACC TAC TAC TGT ACG GGC AAA	643					
	Thr Asn Ala Thr Val Glu Asp Ser Gly Thr Tyr Tyr Cys Thr Gly Lys						
	165 170 175						
40	GTG TGG CAG CTG GAC TAT GAG TCT GAG CCC CTC AAC ATT ACT GTA ATA	691					
	Val Trp Gln Leu Asp Tyr Glu Ser Glu Pro Leu Asn Ile Thr Val Ile						
	180 185 190 195						
	AAA GCT CCG CGT GAG AAG TAC TGG CTA CAA TTT TTT ATC CCA TTG TTG	739					
	Lys Ala Pro Arg Glu Lys Tyr Trp Leu Gln Phe Phe Ile Pro Leu Leu						
	200 205 210						
45	GTG GTG ATT CTG TTT GCT GTG GAC ACA GGA TTA TTT ATC TCA ACT CAG	787					
	Val Val Ile Leu Phe Ala Val Asp Thr Gly Leu Phe Ile Ser Thr Gln						
	215 220 225						
	CAG CAG GTC ACA TTT CTC TTG AAG ATT AAG AGA ACC AGG AAA GGC TTC	835					
	Gln Gln Val Thr Phe Leu Leu Lys Ile Lys Arg Thr Arg Lys Gly Phe						
50	230 235 240						

-37-

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 Arg Leu Leu Asn Pro His Pro Lys Pro Asn Pro Lys Asn Asn
 245 250 255

5 TATAATTACT CAAGAAATAT TTGCAACATT AGTTTTTTTC CAGCATCAGC AATTGCTACT 940
 CAATTGTCAA ACACAGCTTG CAATATACAT AGAAACGTCT GTGCTCAAGG ATTTATAGAA 1000
 ATGCTTCATT AACTTGAGTG AACTGGTTA AGTGGCATGT AATAGTAAGT GCTCAATTAA 1060
 CATTGGTTGA ATAAATGAGA GAATGAATAG ATTCATTIAT TAGCATTTGT AAAAGAGATG 1120
 TTCAATTTCA ATAAATATAA TATAAAACCA TGTAACAGAA TGCTTCTGAG TAAAAAATAA 1180
 AAAAAAAAAA AAAAAAAAAA 1198

10 (2) INFORMATION FOR SEQ ID NO:2:

(i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 257 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear

15 (ii) MOLECULE TYPE: protein

(iii) SEQUENCE DESCRIPTION: SEQ ID NO:2:

Met Ala Pro Ala Met Glu Ser Pro Thr Leu Leu Cys Val Ala Leu Leu
 1 5 10 15
 20 Phe Phe Ala Pro Asp Gly Val Leu Ala Val Pro Gln Lys Pro Lys Val
 20 25 30
 Ser Leu Asn Pro Pro Trp Asn Arg Ile Phe Lys Gly Glu Asn Val Thr
 35 40 45
 Leu Thr Cys Asn Gly Asn Asn Phe Phe Glu Val Ser Ser Thr Lys Trp
 50 55 60
 25 Phe His Asn Gly Ser Leu Ser Glu Glu Thr Asn Ser Ser Leu Asn Ile
 65 70 75 80
 Val Asn Ala Lys Phe Glu Asp Ser Gly Glu Tyr Lys Cys Gln His Gln
 85 90 95
 30 Gln Val Asn Glu Ser Glu Pro Val Tyr Leu Glu Val Phe Ser Asp Trp
 100 105 110
 Leu Leu Leu Gln Ala Ser Ala Glu Val Val Met Glu Gly Gln Pro Leu
 115 120 125
 Phe Leu Arg Cys His Gly Trp Arg Asn Trp Asp Val Tyr Lys Val Ile
 130 135 140
 35 Tyr Tyr Lys Asp Gly Glu Ala Leu Lys Tyr Trp Tyr Glu Asn His Asn
 145 150 155 160
 Ile Ser Ile Thr Asn Ala Thr Val Glu Asp Ser Gly Thr Tyr Tyr Cys
 165 170 175
 40 Thr Gly Lys Val Trp Gln Leu Asp Tyr Glu Ser Glu Pro Leu Asn Ile
 180 185 190
 Thr Val Ile Lys Ala Pro Arg Glu Lys Tyr Trp Leu Gln Phe Phe Ile
 195 200 205
 Pro Leu Leu Val Val Ile Leu Phe Ala Val Asp Thr Gly Leu Phe Ile
 210 215 220
 45 Ser Thr Gln Gln Gln Val Thr Phe Leu Leu Lys Ile Lys Arg Thr Arg
 225 230 235 240

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Lys Gly Phe Arg Leu Leu Asn Pro His Pro Lys Pro Asn Pro Lys Asn
 245 250 255

Asn

(2) INFORMATION FOR SEQ ID NO:3:

- 5 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 1198 nucleotides
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: single
 (D) TOPOLOGY: linear

- 10 (ii) MOLECULE TYPE: cDNA

(iii) SEQUENCE DESCRIPTION: SEQ ID NO:3:

TTTTTTTTTT	TTTTTTTTTT	TTTTTTTACT	CAGAAGCATT	CTGTTACATG	GTTTTATATT	60
TATTTTATTG	AAATTGAACA	TCTCTTTTAC	AAATGCTAAT	AAATGAATCT	ATTCATTCTC	120
TCATTTATTC	AACCAATGTT	AATTGAGCAC	TTACTATTAC	ATGCCACTTA	ACCAGTTTCA	180
15 CTCAGTTTAA	TGAAGCATT	CTATAAATCC	TTGAGCACAG	ACGTTTCTAT	GTATATTGCA	240
AGCTGTGTTT	GACAATTGAG	TAGCAATTGC	TGATGCTGGA	AAAAAACTAA	TGTTGCAAAT	300
ATTTCTTGAG	TAATTATATC	AGTTGTTTTT	GGGGTTTGGC	TTAGGATGTG	GGTTCAGAAG	360
TCTGAAGCCT	TTCCTGGTTC	TCTTAATCTT	CAAGAGAAAT	GTGACCTGCT	GCTGAGTTGA	420
GATAAATAAT	CCTGTGTCCA	CAGCAAACAG	AATCACCACC	AACAATGGGA	TAAAAAATTG	480
20 TAGCCAGTAC	TTCTCACGCG	GAGCTTTTAT	TACAGTAATG	TTGAGGGGCT	CAGACTCATA	540
GTCCAGCTGC	CACACTTTGC	CCGTACAGTA	GTAGGTTCCA	CTGTCTTCAA	CTGTGGCATT	600
TGTAATGGAG	ATGTTGTGGT	TCTCATACCA	GTAAGTTGGA	GCTTCACCAT	CCTTATAATA	660
GATCACCTTG	TACACATCCC	AGTTCCTCCA	ACCATGGCAC	CTGAGGAAGA	GGGGCTGGCC	720
CTCCATCACC	ACCTCAGCAG	AGGCCTGAAG	GAGCAGCCAG	TCACTGAAGA	CTTCCAGGTA	780
25 CACAGGTTCA	CTCTCATTAA	CTTGTTGGTG	CTGACATTTG	TATTCCTCAC	TGTCCTTCAA	840
TTTGGCATT	ACAATATTCA	AACTTGAATT	TGTCTCTTCT	GAAAGGCTGC	CATTGTGGAA	900
CCATTGGGTG	GAACGTGACT	CAAAGAAATT	GTTCCCATT	CATGTAAGAG	TCACATTCTC	960
TCCTTTAAAT	ATTCTATTCC	ATGGAGGGTT	CAAGGAGACC	TTAGGTTTCT	GAGGGACTGC	1020
TAACACGCCA	TCTGGAGCGA	AGAACAGTAA	GGCTACACAC	AGTAGAGTAG	GGGATTCCAT	1080
30 GGCAGGAGCC	ATCTTCTTCA	TGGACTCCTG	GTGCTTACTG	TGCTGGAGAG	ATCTAAGGCT	1140
TCAAATATAG	GCCCATGCTC	GGTGGTAGAC	AGGTGGAGGA	TGCTGGAGAC	TCCTTAGTA	1198

(2) INFORMATION FOR SEQ ID NO:4:

- 35 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 774 nucleotides
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: single
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(iii) FEATURE:

- 40 (A) NAME/KEY: CDS
 (B) LOCATION: 1..774

(iv) SEQUENCE DESCRIPTION: SEQ ID NO:4:

ATG GCT CCT GCC ATG GAA TCC CCT ACT CTA CTG TGT GTA GCC TTA CTG	48
Met Ala Pro Ala Met Glu Ser Pro Thr Leu Leu Cys Val Ala Leu Leu	
1 5 10 15	
TTC TTC GCT CCA GAT GGC GTG TTA GCA GTC CCT CAG AAA CCT AAG GTC	96
Phe Phe Ala Pro Asp Gly Val Leu Ala Val Pro Gln Lys Pro Lys Val	
20 25 30	
TCC TTG AAC CCT CCA TGG AAT AGA ATA TTT AAA GGA GAG AAT GTG ACT	144
Ser Leu Asn Pro Pro Trp Asn Arg Ile Phe Lys Gly Glu Asn Val Thr	
35 40 45	

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	CTT	ACA	TGT	AAT	GGG	AAC	AAT	TTC	TTT	GAA	GTC	AGT	TCC	ACC	AAA	TGG	192
	Leu	Thr	Cys	Asn	Gly	Asn	Asn	Phe	Phe	Glu	Val	Ser	Ser	Thr	Lys	Trp	
		50					55					60					
5	TTC	CAC	AAT	GGC	AGC	CTT	TCA	GAA	GAG	ACA	AAT	TCA	AGT	TTG	AAT	ATT	240
	Phe	His	Asn	Gly	Ser	Leu	Ser	Glu	Glu	Thr	Asn	Ser	Ser	Leu	Asn	Ile	
	65					70					75					80	
	GTG	AAT	GCC	AAA	TTT	GAA	GAC	AGT	GGA	GAA	TAC	AAA	TGT	CAG	CAC	CAA	288
	Val	Asn	Ala	Lys	Phe	Glu	Asp	Ser	Gly	Glu	Tyr	Lys	Cys	Gln	His	Gln	
					85					90					95		
10	CAA	GTT	AAT	GAG	AGT	GAA	CCT	GTG	TAC	CTG	GAA	GTC	TTC	AGT	GAC	TGG	336
	Gln	Val	Asn	Glu	Ser	Glu	Pro	Val	Tyr	Leu	Glu	Val	Phe	Ser	Asp	Trp	
				100					105					110			
15	CTG	CTC	CTT	CAG	GCC	TCT	GCT	GAG	GTG	GTG	ATG	GAG	GGC	CAG	CCC	CTC	384
	Leu	Leu	Leu	Gln	Ala	Ser	Ala	Glu	Val	Val	Met	Glu	Gly	Gln	Pro	Leu	
			115					120					125				
	TTC	CTC	AGG	TGC	CAT	GGT	TGG	AGG	AAC	TGG	GAT	GTG	TAC	AAG	GTG	ATC	432
	Phe	Leu	Arg	Cys	His	Gly	Trp	Arg	Asn	Trp	Asp	Val	Tyr	Lys	Val	Ile	
		130					135					140					
20	TAT	TAT	AAG	GAT	GGT	GAA	GCT	CTC	AAG	TAC	TGG	TAT	GAG	AAC	CAC	AAC	480
	Tyr	Tyr	Lys	Asp	Gly	Glu	Ala	Leu	Lys	Tyr	Trp	Tyr	Glu	Asn	His	Asn	
	145				150						155					160	
	ATC	TCC	ATT	ACA	AAT	GCC	ACA	GTT	GAA	GAC	AGT	GGA	ACC	TAC	TAC	TGT	528
	Ile	Ser	Ile	Thr	Asn	Ala	Thr	Val	Glu	Asp	Ser	Gly	Thr	Tyr	Tyr	Cys	
					165					170					175		
25	ACG	GGC	AAA	GTG	TGG	CAG	CTG	GAC	TAT	GAG	TCT	GAG	CCC	CTC	AAC	ATT	576
	Thr	Gly	Lys	Val	Trp	Gln	Leu	Asp	Tyr	Glu	Ser	Glu	Pro	Leu	Asn	Ile	
				180					185					190			
30	ACT	GTA	ATA	AAA	GCT	CCG	CGT	GAG	AAG	TAC	TGG	CTA	CAA	TTT	TTT	ATC	624
	Thr	Val	Ile	Lys	Ala	Pro	Arg	Glu	Lys	Tyr	Trp	Leu	Gln	Phe	Phe	Ile	
			195					200					205				
	CCA	TTG	TTG	GTG	GTG	ATT	CTG	TTT	GCT	GTG	GAC	ACA	GGA	TTA	TTT	ATC	672
	Pro	Leu	Leu	Val	Val	Ile	Leu	Phe	Ala	Val	Asp	Thr	Gly	Leu	Phe	Ile	
		210					215					220					
35	TCA	ACT	CAG	CAG	CAG	GTC	ACA	TTT	CTC	TTG	AAG	ATT	AAG	AGA	ACC	AGG	720
	Ser	Thr	Gln	Gln	Gln	Val	Thr	Phe	Leu	Leu	Lys	Ile	Lys	Arg	Thr	Arg	
	225					230					235					240	
	AAA	GGC	TTC	AGA	CTT	CTG	AAC	CCA	CAT	CCT	AAG	CCA	AAC	CCC	AAA	AAC	768
	Lys	Gly	Phe	Arg	Leu	Leu	Asn	Pro	His	Pro	Lys	Pro	Asn	Pro	Lys	Asn	
					245					250					255		
40	AAC	TGA															774
	Asn																

(2) INFORMATION FOR SEQ ID NO:5:

- (i) SEQUENCE CHARACTERISTICS:
- (A) LENGTH: 774 nucleotides
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

45

(ii) MOLECULE TYPE: cDNA

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(iii) SEQUENCE DESCRIPTION: SEQ ID NO:5:

	TCAGTTGTTT	TTGGGGTTTG	GCTTAGGATG	TGGGTTTCAGA	AGTCTGAAGC	CTTTCCTGGT	60
	TCTCTTAATC	TTCAAGAGAA	ATGTGACCTG	CTGCTGAGTT	GAGATAAATA	ATCCTGTGTC	120
5	CACAGCAAAC	AGAATCACCA	CCAACAATGG	GATAAAAAAT	TGTAGCCAGT	ACTTCTCAGC	180
	CGGAGCTTTT	ATTACAGTAA	TGTTGAGGGG	CTCAGACTCA	TAGTCCAGCT	GCCACACTTT	240
	GCCCCGTACAG	TAGTAGGTTT	CACTGTCTTC	AACTGTGGCA	TTTGTAATGG	AGATGTTGTG	300
	GTTCTCATAC	CAGTACTTGA	GAGCTTCACC	ATCCTTATAA	TAGATCACCT	TGTACACATC	360
	CCAGTTCCTC	CAACCATGGC	ACCTGAGGAA	GAGGGGCTGG	CCCTCCATCA	CCACCTCAGC	420
	AGAGGCCTGA	AGGAGCAGCC	AGTCACTGAA	GACTTCCAGG	TACACAGGTT	CACTCTCAT	480
10	AACTTGTTGG	TGCTGACATT	TGTATTCTCC	ACTGTCTTCA	AATTTGGCAT	TCACAATATT	540
	CAAACCTGAA	TTTGTCCTTT	CTGAAAGGCT	GCCATTGTGG	AACCATTTGG	TGGAACCTGAC	600
	TTCAAAGAAA	TTGTTCCCAT	TACATGTAAG	AGTCACATTC	TCTCCTTTAA	ATATTCTATT	660
	CCATGGAGGG	TTCAAGGAGA	CCTTAGGTTT	CTGAGGGACT	GCTAACACGC	CATCTGGAGC	720
	GAAGAACAGT	AAGGCTACAC	ACAGTAGAGT	AGGGGATTCC	ATGGCAGGAG	CCAT	774

15 (2) INFORMATION FOR SEQ ID NO:6:

(i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 232 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear

20 (ii) MOLECULE TYPE: protein

(iii) SEQUENCE DESCRIPTION: SEQ ID NO:6:

	Val	Pro	Gln	Lys	Pro	Lys	Val	Ser	Leu	Asn	Pro	Pro	Trp	Asn	Arg	Ile	
	1				5					10					15		
25	Phe	Lys	Gly	Glu	Asn	Val	Thr	Leu	Thr	Cys	Asn	Gly	Asn	Asn	Phe	Phe	
				20					25					30			
	Glu	Val	Ser	Ser	Thr	Lys	Trp	Phe	His	Asn	Gly	Ser	Leu	Ser	Glu	Glu	
			35					40					45				
	Thr	Asn	Ser	Ser	Leu	Asn	Ile	Val	Asn	Ala	Lys	Phe	Glu	Asp	Ser	Gly	
		50				55						60					
30	Glu	Tyr	Lys	Cys	Gln	His	Gln	Gln	Val	Asn	Glu	Ser	Glu	Pro	Val	Tyr	
		65			70					75					80		
	Leu	Glu	Val	Phe	Ser	Asp	Trp	Leu	Leu	Leu	Gln	Ala	Ser	Ala	Glu	Val	
				85					90						95		
35	Val	Met	Glu	Gly	Gln	Pro	Leu	Phe	Leu	Arg	Cys	His	Gly	Trp	Arg	Asn	
			100					105					110				
	Trp	Asp	Val	Tyr	Lys	Val	Ile	Tyr	Tyr	Lys	Asp	Gly	Glu	Ala	Leu	Lys	
		115						120				125					
	Tyr	Trp	Tyr	Glu	Asn	His	Asn	Ile	Ser	Ile	Thr	Asn	Ala	Thr	Val	Glu	
		130				135						140					
40	Asp	Ser	Glv	Thr	Tyr	Tyr	Cys	Thr	Gly	Lys	Val	Trp	Gln	Leu	Asp	Tyr	
		145				150					155				160		
	Glu	Ser	Glu	Pro	Leu	Asn	Ile	Thr	Val	Ile	Lys	Ala	Pro	Arg	Glu	Lys	
				165						170					175		
45	Tyr	Trp	Leu	Gln	Phe	Phe	Ile	Pro	Leu	Leu	Val	Val	Ile	Leu	Phe	Ala	
			180					185						190			
	Val	Asp	Thr	Gly	Leu	Phe	Ile	Ser	Thr	Gln	Gln	Gln	Val	Thr	Phe	Leu	
		195						200					205				

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Leu Lys Ile Lys Arg Thr Arg Lys Gly Phe Arg Leu Leu Asn Pro His
 210 215 220

Pro Lys Pro Asn Pro Lys Asn Asn
 225 230

5 (2) INFORMATION FOR SEQ ID NO:7:

(i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 699 nucleotides
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: single
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(iii) FEATURE:
 (A) NAME/KEY: CDS
 (B) LOCATION: 1..699

15 (iii) SEQUENCE DESCRIPTION: SEQ ID NO:7:

	GTC CCT CAG AAA CCT AAG GTC TCC TTG AAC CCT CCA TGG AAT AGA ATA	48
	Val Pro Gln Lys Pro Lys Val Ser Leu Asn Pro Pro Trp Asn Arg Ile	
	1 5 10 15	
20	TTT AAA GGA GAG AAT GTG ACT CTT ACA TGT AAT GGG AAC AAT TTC TTT	96
	Lys Gly Glu Asn Val Thr Leu Thr Cys Phe Asn Gly Asn Asn Phe Phe	
	20 25 30	
	GAA GTC AGT TCC ACC AAA TGG TTC CAC AAT GGC AGC CTT TCA GAA GAG	144
	Glu Val Ser Ser Thr Lys Trp Phe His Asn Gly Ser Leu Ser Glu Glu	
	35 40 45	
25	ACA AAT TCA AGT TTG AAT ATT GTG AAT GCC AAA TTT GAA GAC AGT GGA	192
	Thr Asn Ser Ser Leu Asn Ile Val Asn Ala Lys Phe Glu Asp Ser Gly	
	50 55 60	
	GAA TAC AAA TGT CAG CAC CAA CAA GTT AAT GAG AGT GAA CCT GTG TAC	240
30	Glu Tyr Lys Cys Gln His Gln Gln Val Asn Glu Ser Glu Pro Val Tyr	
	65 70 75 80	
	CTG GAA GTC TTC AGT GAC TGG CTG CTC CTT CAG GCC TCT GCT GAG GTG	288
	Leu Glu Val Phe Ser Asp Trp Leu Leu Leu Gln Ala Ser Ala Glu Val	
	85 90 95	
35	GTG ATG GAG GGC CAG CCC CTC TTC CTC AGG TGC CAT GGT TGG AGG AAC	336
	Val Met Glu Gly Gln Pro Leu Phe Leu Arg Cys His Gly Trp Arg Asn	
	100 105 110	
	TGG GAT GTG TAC AAG GTG ATC TAT TAT AAG GAT GGT GAA GCT CTC AAG	384
	Trp Asp Val Tyr Lys Val Ile Tyr Tyr Lys Asp Gly Glu Ala Leu Lys	
	115 120 125	
40	TAC TGG TAT GAG AAC CAC AAC ATC TCC ATT ACA AAT GCC ACA GTT GAA	432
	Tyr Trp Tyr Glu Asn His Asn Ile Ser Ile Thr Asn Ala Thr Val Glu	
	130 135 140	
	GAC AGT GGA ACC TAC TAC TGT ACG GGC AAA GTG TGG CAG CTG GAC TAT	480
45	Asp Ser Gly Thr Tyr Tyr Cys Thr Gly Lys Val Trp Gln Leu Asp Tyr	
	145 150 155 160	
	GAG TCT GAG CCC CTC AAC ATT ACT GTA ATA AAA GCT CCG CGT GAG AAG	528
	Glu Ser Glu Pro Leu Asn Ile Thr Val Ile Lys Ala Pro Arg Glu Lys	
	165 170 175	

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	TAC Tyr	TGG Trp	CTA Leu	CAA Gln 180	TTT Phe	TTT Phe	ATC Ile	CCA Pro	TTG Leu 185	TTG Leu	GTG Val	GTG Val	ATT Ile	CTG Leu 190	TTT Phe	GCT Ala	576
5	GTG Val	GAC Asp	ACA Thr 195	GGA Gly	TTA Leu	TTT Phe	ATC Ile	TCA Ser 200	ACT Thr	CAG Gln	CAG Gln	CAG Gln	GTC Val 205	ACA Thr	TTT Phe	CTC Leu	624
	TTG Leu	AAG Lys 210	ATT Ile	AAG Lys	AGA Arg	ACC Thr	AGG Arg 215	AAA Lys	GGC Gly	TTC Phe	AGA Arg	CTT Leu 220	CTG Leu	AAC Asn	CCA Pro	CAT His	672
10	CCT Pro 225	AAG Lys	CCA Pro	AAC Asn	CCC Pro	AAA Lys 230	AAC Asn	AAC Asn	TGA								699

(2) INFORMATION FOR SEQ ID NO:8:

15 (i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 32 bases
(B) TYPE: nucleic acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: primer

20 (iii) SEQUENCE DESCRIPTION: SEQ ID NO:8:

CGCGGATCCT ATAAATATGG CTCCTGCCAT GG

(2) INFORMATION FOR SEQ ID NO:9:

25 (i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 26 bases
(B) TYPE: nucleic acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: primer

(iii) SEQUENCE DESCRIPTION: SEQ ID NO:9:

30 GCGGAATTCT TAAGCTTTTA TTACAG

(2) INFORMATION FOR SEQ ID NO:10:

(i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 591 nucleotides
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: single
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

40 (iii) FEATURE:
(A) NAME/KEY: CDS
(B) LOCATION: 1..591

(iv) SEQUENCE DESCRIPTION: SEQ ID NO:10:

ATG GCT CCT GCC ATG GAA TCC CCT ACT CTA CTG TGT GTA GCC TTA CTG 48
Met Ala Pro Ala Met Glu Ser Pro Thr Leu Leu Cys Val Ala Leu Leu
1 5 10 15

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	TTC	TTC	GCT	CCA	GAT	GGC	GTG	TTA	GCA	GTC	CCT	CAG	AAA	CCT	AAG	GTC	96
	Phe	Phe	Ala	Pro	Asp	Gly	Val	Leu	Ala	Val	Pro	Gln	Lys	Pro	Lys	Val	
				20					25					30			
5	TCC	TTG	AAC	CCT	CCA	TGG	AAT	AGA	ATA	TTT	AAA	GGA	GAG	AAT	GTG	ACT	144
	Ser	Leu	Asn	Pro	Pro	Trp	Asn	Arg	Ile	Phe	Lys	Gly	Glu	Asn	Val	Thr	
			35				40					45					
	CTT	ACA	TGT	AAT	GGG	AAC	AAT	TTC	TTT	GAA	GTC	AGT	TCC	ACC	AAA	TGG	192
	Leu	Thr	Cys	Asn	Gly	Asn	Asn	Phe	Phe	Glu	Val	Ser	Ser	Thr	Lys	Trp	
		50				55						60					
10	TTC	CAC	AAT	GGC	AGC	CTT	TCA	GAA	GAG	ACA	AAT	TCA	AGT	TTG	AAT	ATT	240
	Phe	His	Asn	Gly	Ser	Leu	Ser	Glu	Glu	Thr	Asn	Ser	Ser	Leu	Asn	Ile	
	65					70				75						80	
15	GTG	AAT	GCC	AAA	TTT	GAA	GAC	AGT	GGA	GAA	TAC	AAA	TGT	CAG	CAC	CAA	288
	Val	Asn	Ala	Lys	Phe	Glu	Asp	Ser	Gly		Tyr	Lys	Cys	Gln	His	Gln	
					85					90					95		
	CAA	GTT	AAT	GAG	AGT	GAA	CCT	GTG	TAC	CTG	GAA	GTC	TTC	AGT	GAC	TGG	336
	Gln	Val	Asn	Glu	Ser	Glu	Pro	Val	Tyr	Leu	Glu	Val	Phe	Ser	Asp	Trp	
				100					105					110			
20	CTG	CTC	CTT	CAG	GCC	TCT	GCT	GAG	GTG	GTG	ATG	GAG	GGC	CAG	CCC	CTC	384
	Leu	Leu	Leu	Gln	Ala	Ser	Ala	Glu	Val	Val	Met	Glu	Gly	Gln	Pro	Leu	
			115					120					125				
	TTC	CTC	AGG	TGC	CAT	GGT	TGG	AGG	AAC	TGG	GAT	GTG	TAC	AAG	GTG	ATC	432
	Phe	Leu	Arg	Cys	His	Gly	Trp	Arg	Asn	Trp	Asp	Val	Tyr	Lys	Val	Ile	
		130				135						140					
25	TAT	TAT	AAG	GAT	GGT	GAA	GCT	CTC	AAG	TAC	TGG	TAT	GAG	AAC	CAC	AAC	480
	Tyr	Tyr	Lys	Asp	Gly	Glu	Ala	Leu	Lys	Tyr	Trp	Tyr	Glu	Asn	His	Asn	
	145					150					155					160	
30	ATC	TCC	ATT	ACA	AAT	GCC	ACA	GTT	GAA	GAC	AGT	GGA	ACC	TAC	TAC	TGT	528
	Ile	Ser	Ile	Thr	Asn	Ala	Thr	Val	Glu	Asp	Ser	Gly	Thr	Tyr	Tyr	Cys	
					165					170					175		
	ACG	GGC	AAA	GTG	TGG	CAG	CTG	GAC	TAT	GAG	TCT	GAG	CCC	CTC	AAC	ATT	576
	Thr	Gly	Lys	Val	Trp	Gln	Leu	Asp	Tyr	Glu	Ser	Glu	Pro	Leu	Asn	Ile	
				180					185					190			
35	ACT	GTA	ATA	AAA	GCT												591
	Thr	Val	Ile	Lys	Ala												
				195													

(2) INFORMATION FOR SEQ ID NO:11:

- (i) SEQUENCE CHARACTERISTICS:
- (A) LENGTH: 197 amino acids
- (B) TYPE: amino acid
- (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: protein
- (iii) SEQUENCE DESCRIPTION: SEQ ID NO:11:
- | | | | | | | | | | | | | | | | | |
|----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|
| 45 | Met | Ala | Pro | Ala | Met | Glu | Ser | Pro | Thr | Leu | Leu | Cys | Val | Ala | Leu | Leu |
| | 1 | | | | 5 | | | | | 10 | | | | | 15 | |
| | Phe | Phe | Ala | Pro | Asp | Gly | Val | Leu | Ala | Val | Pro | Gln | Lys | Pro | Lys | Val |
| | | | | 20 | | | | | 25 | | | | | 30 | | |

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Ser Leu Asn Pro Pro Trp Asn Arg Ile Phe Lys Gly Glu Asn Val Thr
 35 40 45
 Leu Thr Cys Asn Gly Asn Asn Phe Phe Glu Val Ser Ser Thr Lys Trp
 50 55 60
 5 Phe His Asn Gly Ser Leu Ser Glu Glu Thr Asn Ser Ser Leu Asn Ile
 65 70 75 80
 Val Asn Ala Lys Phe Glu Asp Ser Gly Glu Tyr Lys Cys Gln His Gln
 85 90 95
 10 Gln Val Asn Glu Ser Glu Pro Val Tyr Leu Glu Val Phe Ser Asp Trp
 100 105 110
 Leu Leu Leu Gln Ala Ser Ala Glu Val Val Met Glu Gly Gln Pro Leu
 115 120 125
 Phe Leu Arg Cys His Gly Trp Arg Asn Trp Asp Val Tyr Lys Val Ile
 130 135 140
 15 Tyr Tyr Lys Asp Gly Glu Ala Leu Lys Tyr Trp Tyr Glu Asn His Asn
 145 150 155 160
 Ile Ser Ile Thr Asn Ala Thr Val Glu Asp Ser Gly Thr Tyr Tyr Cys
 165 170 175
 20 Thr Gly Lys Val Trp Gln Leu Asp Tyr Glu Ser Glu Pro Leu Asn Ile
 180 185 190
 Thr Val Ile Lys Ala
 195

(2) INFORMATION FOR SEQ ID NO:12:

- (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 516 nucleotides
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: single
 (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: cDNA
- 30 (iii) FEATURE:
 (A) NAME/KEY: CDS
 (B) LOCATION: 1..516
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:12:

35	GTC CCT GAG AAA CCT AAG GTC TCC TTG AAC CCT CCA TGG AAT AGA ATA	48
	Val Pro Gln Lys Pro Lys Val Ser Leu Asn Pro Pro Trp Asn Arg Ile	
	1 5 10 15	
	TTT AAA GGA GAG AAT GTG ACT CTT ACA TGT AAT GGG AAC AAT TTC TTT	96
	Phe Lys Gly Glu Asn Val Thr Leu Thr Cys Asn Gly Asn Asn Phe Phe	
	20 25 30	
40	GAA GTC AGT TCC ACC AAA TGG TTC CAC AAT GGC AGC CTT TCA GAA GAG	144
	Glu Val Ser Ser Thr Lys Trp Phe His Asn Gly Ser Leu Ser Glu Glu	
	35 40 45	
45	ACA AAT TCA AGT TTG AAT ATT GTG AAT GCC AAA TTT GAA GAC AGT GGA	192
	Thr Asn Ser Ser Leu Asn Ile Val Asn Ala Lys Phe Glu Asp Ser Gly	
	50 55 60	

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Asp	Ser	Gly	Thr	Tyr	Tyr	Cys	Thr	Gly	Lys	Val	Trp	Gln	Leu	Asp	Tyr
145					150					155					160
Glu	Ser	Glu	Pro	Leu	Asn	Ile	Thr	Val	Ile	Lys	Ala				
				165					170						

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While various embodiments of the present invention have been described in detail, it is apparent that modifications and adaptations of those embodiments will occur to those skilled in the art. It is to be expressly understood, however, that such modifications and adaptations are within the scope of the present invention, as set forth

5 in the following claims.

What is claimed is:

1. A method to detect IgE comprising:
 - (a) contacting an isolated human $\text{Fc}_\epsilon\text{R}$ receptor ($\text{Fc}_\epsilon\text{R}$) molecule with a putative IgE-containing composition under conditions suitable for formation of a $\text{Fc}_\epsilon\text{R}$ molecule:IgE complex, wherein said IgE is selected from the group consisting of canine IgE, feline IgE and equine IgE; and
 - (b) determining the presence of IgE by detecting said $\text{Fc}_\epsilon\text{R}$ molecule:IgE complex, the presence of said $\text{Fc}_\epsilon\text{R}$ molecule:IgE complex indicating the presence of IgE.
2. A method to detect IgE comprising:
 - (a) contacting a recombinant cell with a putative IgE-containing composition under conditions suitable for formation of a recombinant cell:IgE complex, wherein said recombinant cell is selected from the group consisting of: a recombinant cell expressing a human $\text{Fc}_\epsilon\text{R}$ molecule; and a recombinant cell expressing an antibody that binds selectively to an IgE selected from the group consisting of canine IgE, feline IgE and equine IgE; and
 - (b) determining the presence of IgE by detecting said recombinant cell:IgE complex, the presence of said recombinant cell:IgE complex indicating the presence of IgE.
3. A kit for detecting IgE comprising a human $\text{Fc}_\epsilon\text{R}$ receptor ($\text{Fc}_\epsilon\text{R}$) molecule and a means for detecting an IgE selected from the group consisting of canine IgE, feline IgE and equine IgE.
4. A general allergen kit comprising an allergen common to all regions of the United States and a human $\text{Fc}_\epsilon\text{R}$ receptor ($\text{Fc}_\epsilon\text{R}$) molecule.
5. A method to detect flea allergy dermatitis comprising:
 - (a) immobilizing a flea allergen on a substrate;
 - (b) contacting said flea allergen with a putative IgE-containing composition under conditions suitable for formation of an allergen:IgE complex bound to said substrate;
 - (c) removing non-bound material from said substrate under conditions that retain allergen:IgE complex binding to said substrate; and

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(d) determining the presence of said allergen:IgE complex by contacting said allergen:IgE complex with a $Fc_\epsilon R$ molecule.

6. A kit for detecting flea allergy dermatitis comprising a human Fc_ϵ receptor ($Fc_\epsilon R$) molecule and a flea allergen.

5 7. An isolated human Fc_ϵ receptor ($Fc_\epsilon R$) alpha chain protein, wherein a carbohydrate group of said $Fc_\epsilon R$ alpha chain protein is conjugated to biotin.

8. The invention of Claim 1, 2, 3, 4, 5, 6 or 7, wherein said $Fc_\epsilon R$ molecule comprises at least a portion of a $Fc_\epsilon R$ alpha chain that binds to IgE.

9. The invention of Claim 1, 3, 4, 5, 6 or 7, wherein said $Fc_\epsilon R$ molecule
10 comprises a protein selected from the group consisting of $PhFc_\epsilon R\alpha_{257}$, $PhFc_\epsilon R\alpha_{197}$, $PhFc_\epsilon R\alpha_{232}$ and $PhFc_\epsilon R\alpha_{172}$.

10. The invention of Claim 1, 3, 4, 5, 6 or 7, wherein said $Fc_\epsilon R$ molecule is encoded by a nucleic acid molecule selected from the group consisting of $nhFc_\epsilon R\alpha_{774}$, $nhFc_\epsilon R\alpha_{1198}$, $nhFc_\epsilon R\alpha_{612}$, $nhFc_\epsilon R\alpha_{591}$, $nhFc_\epsilon R\alpha_{699}$ and $nhFc_\epsilon R\alpha_{516}$.

11. The invention of Claim 1, 3, 4, 5, 6 or 7, wherein said $Fc_\epsilon R$ molecule is
15 encoded by a nucleic acid molecule selected from the group consisting of a nucleic acid molecule comprising a nucleic acid sequence selected from the group consisting of SEQ ID NO:1, SEQ ID NO:4, SEQ ID NO:7, SEQ ID NO:10 and SEQ ID NO:12, and a nucleic acid molecule comprising an allelic variant of a nucleic acid molecule
20 comprising any of said nucleic acid sequences.

12. The invention of Claim 1, 3, 4, 5 or 6, wherein said $Fc_\epsilon R$ molecule is conjugated to a detectable marker.

13. The invention of Claim 1, 3, 4, 5 or 6, wherein said $Fc_\epsilon R$ molecule is
25 conjugated to a detectable marker selected from the group consisting of a radioactive label, a fluorescent label, a chemiluminescent label, a chromophoric label and a ligand.

14. The invention of Claim 1, 3, 4, 5 or 6, wherein a carbohydrate group of said $Fc_\epsilon R$ molecule is conjugated to biotin.

15. The method of Claim 1, 2 or 5, wherein said putative IgE-containing composition comprises a composition selected from the group consisting of blood,
30 serum, plasma, urine, tears, aqueous humor, central nervous system fluid (CSF), saliva, lymph, nasal secretions, milk and feces.

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16. The method of Claim 1, 2 or 5, wherein said putative IgE-containing composition comprises serum.
17. The method of Claim 1, 2 or 5, wherein said putative IgE-containing composition comprises a cell that produces IgE.
- 5 18. The method of Claim 1, 2 or 5, wherein said putative IgE-containing composition comprises a cell selected from the group consisting of a myeloma cell and a basophil cell.
19. The method of Claim 1, further comprising the step selected from the group consisting of binding said $Fc_\epsilon R$ molecule to a substrate prior to performing step
10 (a) to form a $Fc_\epsilon R$ molecule-coated substrate; and binding said putative IgE-containing composition to a substrate prior to performing step (a) to form a putative IgE-containing composition-coated substrate, wherein said substrate is selected from the group consisting of a non-coated substrate, a $Fc_\epsilon R$ molecule-coated substrate, an antigen-coated substrate and an anti-IgE antibody-coated substrate.
- 15 20. The method of Claim 19, wherein said antigen is selected from the group consisting of an allergen and a parasitic antigen.
21. The method of Claim 19, further comprising removing non-bound material from said antigen-coated substrate or said antibody-coated substrate under conditions that retain antigen or antibody binding to said substrate.
- 20 22. The method of Claim 5 or 19, wherein said substrate comprises a material selected from the group consisting of plastic, glass, gel, celluloid, paper and particulate material.
23. The method of Claim 1, 2 or 5, wherein said step of determining comprises performing assays selected from the group consisting of enzyme-linked
25 immunoassays, radioimmunoassays, immunoprecipitations, fluorescence immunoassays, chemiluminescent assay, immunoblot assays, lateral flow assays, agglutination assays and particulate-based assays.
24. The method of Claim 1, wherein said step of determining comprises:
(a) contacting said $Fc_\epsilon R$ molecule:IgE complex with an indicator
30 molecule that binds selectively to said $Fc_\epsilon R$ molecule:IgE complex;

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(b) removing substantially all of said indicator molecule that does not selectively bind to Fc_eR molecule:IgE complex; and

(c) detecting said indicator molecule, wherein the presence of said indicator molecule is indicative of the presence of IgE.

5 25. The method of Claim 24, wherein said indicator molecule comprises a compound selected from the group consisting of a Fc_eR molecule, an antigen, an antibody and a lectin.

26. The method of Claim 1, said method comprising the steps of:

10 (a) immobilizing said Fc_eR molecule on a substrate;
(b) contacting said Fc_eR molecule with said putative IgE-containing composition under conditions suitable for formation of an Fc_eR molecule:IgE complex bound to said substrate;

(c) removing non-bound material from said substrate under conditions that retain Fc_eR molecule:IgE complex binding to said substrate; and

15 (d) determining the presence of said Fc_eR molecule:IgE complex.

27. The method of Claim 26, wherein the presence of said Fc_eR molecule:IgE complex is detected by contacting said Fc_eR molecule:IgE complex with a compound selected from the group consisting of an antigen and an antibody that binds selectively to IgE.

20 28. The method of Claim 27, wherein said compound comprises a detectable marker.

29. The method of Claim 1, said method comprising the steps of:

25 (a) immobilizing a desired antigen on a substrate;
(b) contacting said antigen with said putative IgE-containing composition under conditions suitable for formation of an antigen:IgE complex bound to said substrate;

(c) removing non-bound material from said substrate under conditions that retain antigen:IgE complex binding to said substrate; and

30 (d) determining the presence of said antigen:IgE complex by contacting said antigen:IgE complex with said Fc_eR molecule.

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30. The method of Claim 1, said method comprising the steps of:
- (a) immobilizing an antibody that binds selectively to IgE on a substrate;
 - (b) contacting said antibody with said putative IgE-containing composition under conditions suitable for formation of an antibody:IgE complex bound to said substrate;
 - (c) removing non-bound material from said substrate under conditions that retain antibody:IgE complex binding to said substrate; and
 - (d) determining the presence of said antibody:IgE complex by contacting said antibody:IgE complex with said Fc_εR molecule.
31. The method of Claim 1, said method comprising the steps of:
- (a) immobilizing said putative IgE-containing composition on a substrate;
 - (b) contacting said composition with said Fc_εR molecule under conditions suitable for formation of an Fc_εR molecule:IgE complex bound to said substrate;
 - (c) removing non-bound material from said substrate under conditions that retain Fc_εR molecule:IgE complex binding to said substrate; and
 - (d) determining the presence of said Fc_εR molecule:IgE complex.
32. The invention of Claim 1, 3, 4, 5, 6, 29, 30 or 31, wherein said Fc_εR molecule is conjugated to a detectable marker selected from the group consisting of fluorescein, a radioisotope, a phosphatase, biotin, biotin-related compounds, avidin, avidin-related compounds and a peroxidase.
33. The method of Claim 32, wherein the presence of said Fc_εR molecule:IgE complex is determined by contacting said Fc_εR molecule:IgE complex with an indicator molecule selected from the group consisting of an antibody, an antigen and a lectin.
34. The method of Claim 32, wherein said Fc_εR molecule comprises a detectable marker.
35. The method of Claim 1, wherein said putative IgE-containing composition is obtained from an animal, wherein said animal is selected from the group consisting of a dog and a cat.

36. The method of Claim 1, wherein said method is performed in solution.
37. The method of Claim 2, wherein said recombinant cell expresses a $Fc_\epsilon R$ molecule comprising a protein selected from the group consisting of $PhFc_\epsilon R\alpha_{257}$ and $PhFc_\epsilon R\alpha_{232}$.
- 5 38. The method of Claim 2, wherein said recombinant cell expresses a $Fc_\epsilon R$ molecule encoded by a nucleic acid molecule selected from the group consisting of $nhFc_\epsilon R\alpha_{612}$, $nhFc_\epsilon R\alpha_{591}$, $nhFc_\epsilon R\alpha_{699}$ and $nhFc_\epsilon R\alpha_{516}$.
39. The method of Claim 2, wherein said recombinant cell expresses a $Fc_\epsilon R$ molecule encoded by a nucleic acid molecule selected from the group consisting of a
10 nucleic acid molecule comprising a nucleic acid sequence selected from the group consisting of SEQ ID NO:1 and SEQ ID NO:4, and a nucleic acid molecule comprising an allelic variant of a nucleic acid molecule comprising SEQ ID NO:1 and SEQ ID NO:4.
40. The method of Claim 2, wherein said recombinant cell is a RBL- $hFc_\epsilon R$
15 cell.
41. The kit of Claim 3, wherein said detection means further comprises an antigen selected from the group consisting of an allergen and a parasite antigen, wherein said antigen induces IgE antibody production in animals selected from the group consisting of canines, felines and equines.
- 20 42. The kit of Claim 3, wherein said detection means comprises an antibody that selectively binds to an IgE.
43. The kit of Claim 3, wherein said detection means detects said $Fc_\epsilon R$ molecule.
44. The kit of Claim 3, wherein said $Fc_\epsilon R$ molecule is on the surface of a
25 recombinant cell that expresses said $Fc_\epsilon R$ molecule.
45. The kit of Claim 41, wherein said antigen is immobilized on a substrate.
46. The kit of Claim 45, wherein said substrate comprises a material selected from the group consisting of plastic, glass, gel, celluloid, paper, magnetic resin, poly-vinylidene-fluoride, nylon, nitrocellulose and particulate material.

47. The invention of Claim 5, 19 or 45, wherein said substrate material is selected from the group consisting of latex, polystyrene, nylon, nitrocellulose, agarose and magnetic resin.

48. The invention of Claim 19 or 45, wherein said substrate comprises a
5 shape selected from the group consisting of a well, a plate, a dipstick, a bead, a lateral flow apparatus, a membrane, a filter, a tube, a dish, a celluloid-type matrix and a magnetic particle.

49. The invention of Claim 5, 19 or 45, wherein said substrate comprises an ELISA plate, a dipstick, a radioimmunoassay plate, agarose beads, plastic beads, latex
10 beads, immunoblot membranes and immunoblot papers.

50. The kit of Claim 45, wherein said substrate is latex beads.

51. The kit of Claim 41, wherein said allergen is derived from material selected from the group consisting of fungi, trees, weeds, shrubs, grasses, wheat, corn, soybean, rice, eggs, milk, cheese, bovine, poultry, swine, sheep, yeast, fleas, flies,
15 mosquitos, mites, midges, biting gnats, lice, bees, wasps, ants, true bugs and ticks.

52. The invention of Claim 5 or 51, wherein said flea allergen is a flea saliva antigen.

53. The kit of Claim 3, wherein said parasite antigen is a heartworm antigen.

54. The kit of Claim 3, further comprising an apparatus comprising:
20 (a) a support structure defining a flow path;
(b) a labeling reagent comprising a bead conjugated to said antigen, wherein said labeling reagent is impregnated within the support structure in a labeling zone; and

(c) a capture reagent comprising said Fc_R molecule, wherein said
25 capture reagent is located downstream of said labeling reagent within a capture zone fluidly connected to said labeling zone in such a manner that said labeling reagent can flow from said labeling zone into said capture zone.

55. The kit of Claim 54, wherein said apparatus further comprises a sample receiving zone located along said flow path.

30 56. The kit of Claim 54, wherein said apparatus further comprises an absorbent located at the end of said flow path.

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57. The kit of Claim 55, wherein said sample receiving zone is located upstream of said labeling reagent.

58. The kit of Claim 54, wherein said support structure comprises a material that does not impede the flow of said bead from said labeling zone to said capture zone.

5 59. The kit of Claim 54, wherein said support structure comprises an ionic material.

60. The kit of Claim 54, wherein said support structure comprises a material selected from the group consisting of nitrocellulose, PVDF and carboxymethylcellulose.

61. The kit of Claim 54, wherein said bead comprises a latex bead.

10 62. The kit of Claim 54, wherein said labeling reagent is dried within said labeling zone and said capture reagent is dried within said capture zone.

63. The kit of Claim 4, wherein said allergen is selected from the group consisting of grass, Meadow Fescue, Curly Dock, plantain, Mexican Firebush, Lamb's Quarters, pigweed, ragweed, sage, elm, cocklebur, Box Elder, walnut, cottonwood, ash,
15 birch, cedar, oak, mulberry, cockroach, *Dermataphagoides*, *Alternaria*, *Aspergillus*, *Cladosporium*, *Fusarium*, *Helminthosporium*, *Mucor*, *Penicillium*, *Pullularia*, *Rhizopus* and *Tricophyton*.

64. The kit of Claim 4, wherein said allergen is selected from the group consisting of Johnson Grass, Kentucky Blue Grass, Meadow Fescue, Orchard Grass,
20 Perennial Rye Grass, Redtop Grass, Timothy Grass, Bermuda Grass, Brome Grass, Curly Dock, English Plantain, Mexican Firebush, Lamb's Quarters, Rough Pigweed Short Ragweed, Wormwood Sage, American Elm, Common Cocklebur, Box Elder, Black Walnut, Eastern Cottonwood, Green Ash, River Birch, Red Cedar, Red Oak, Red Mulberry, Cockroach, *Dermataphagoides farinae*, *Alternaria alternata*, *Aspergillus*
25 *fumigatus*, *Cladosporium herbarum*, *Fusarium vasinfectum*, *Helminthosporium sativum*, *Mucor recemosus*, *Penicillium notatum*, *Pullularia pullulans*, *Rhizopus nigricans* and *Tricophyton* spp.

65. The kit of Claim 4, wherein said kit comprises one or more compositions, each composition comprising one allergen.

30 66. The kit of Claim 4, wherein allergen is immobilized to said substrate.

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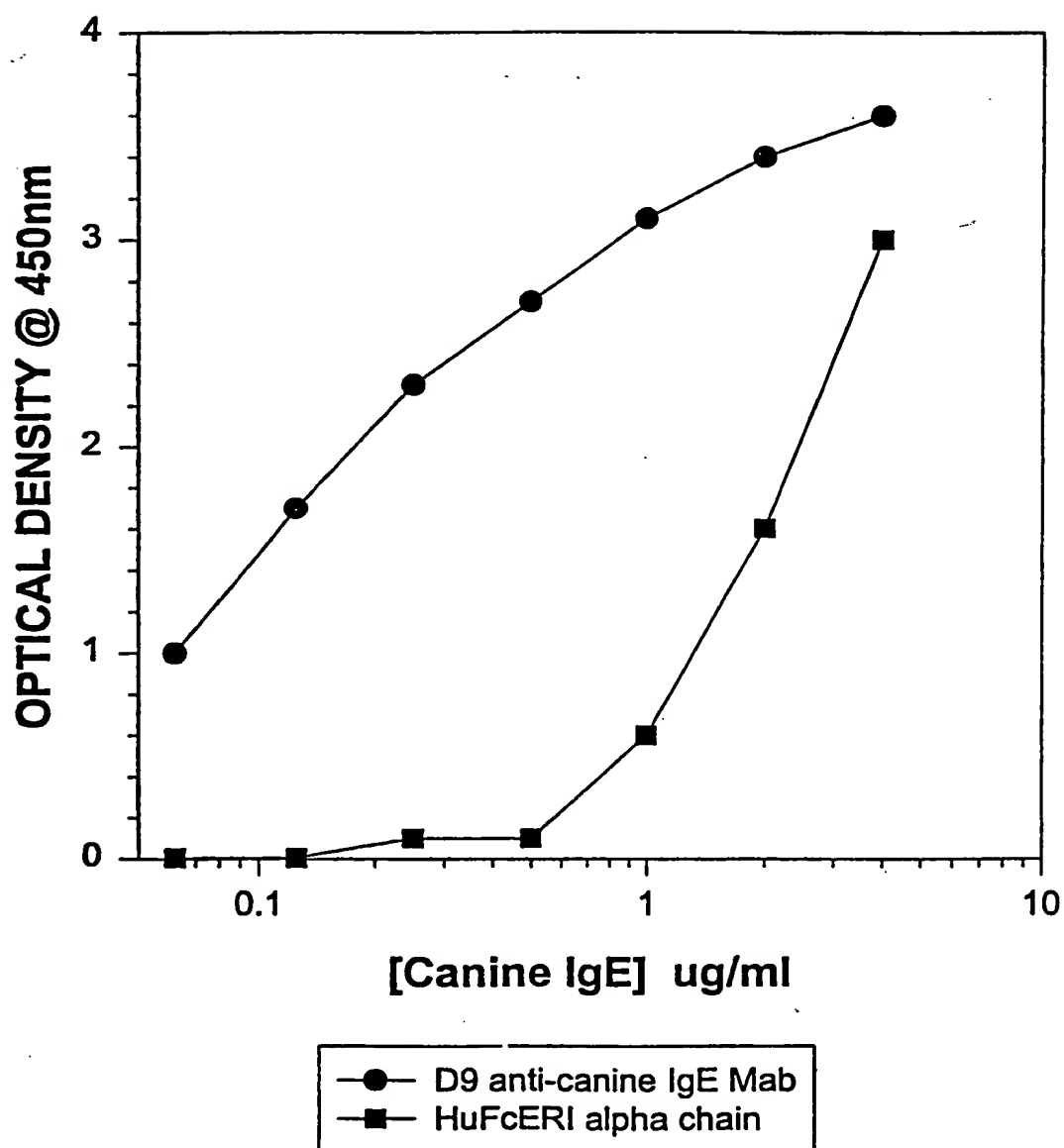
67. The invention of Claim 5 or 6, wherein said flea allergen is selected from the group consisting of flea saliva products and flea saliva proteins.

68. The Fc_εR alpha chain protein of Claim 7, wherein said Fc_εR alpha chain protein is encoded by a nucleic acid molecule selected from the group consisting of a
5 nucleic acid molecule comprising a nucleic acid sequence selected from the group consisting of SEQ ID NO:1, SEQ ID NO:4, SEQ ID NO:7, SEQ ID NO:10 and SEQ ID NO:12, and a nucleic acid molecule comprising an allelic variant of a nucleic acid molecule comprising any of said nucleic acid sequences.

69. The Fc_εR alpha chain protein of Claim 7, wherein said Fc_εR alpha chain
10 protein comprises PhFc_εRα₁₇₂-BIOT.

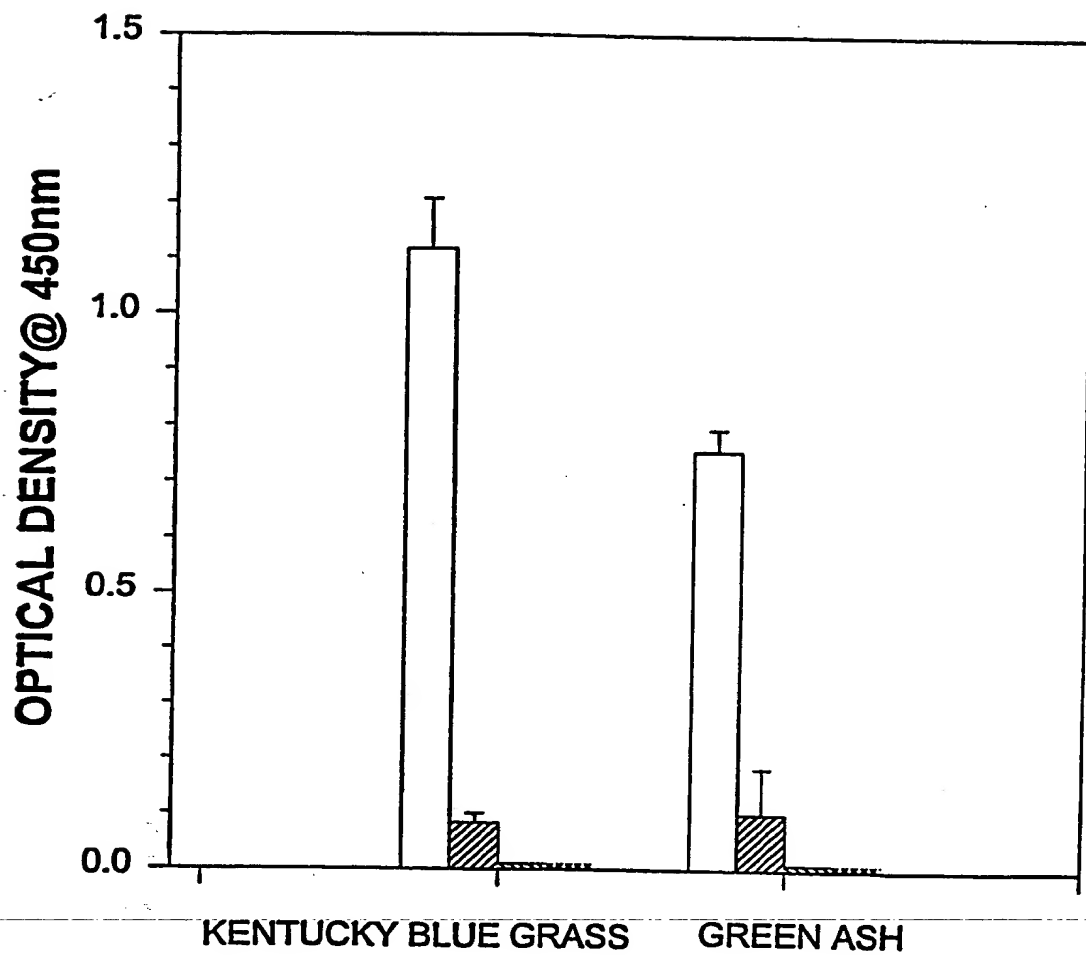
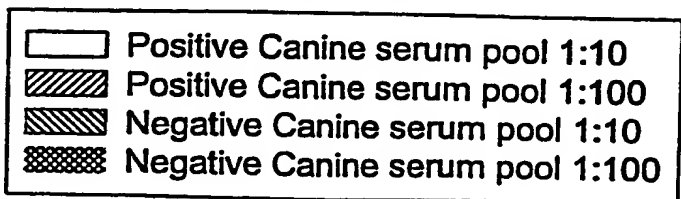
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Fig. 1



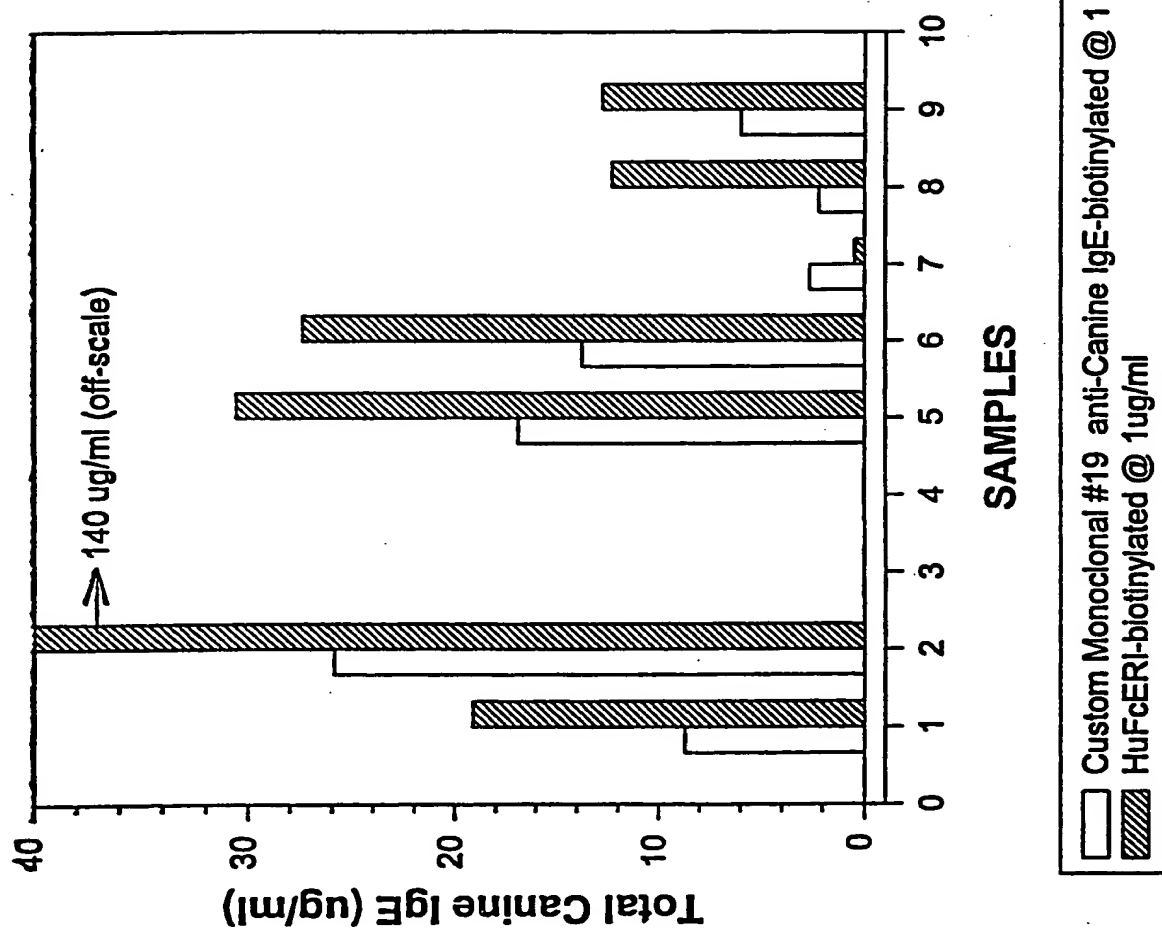
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Fig. 2

**SAMPLES**

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Fig. 3



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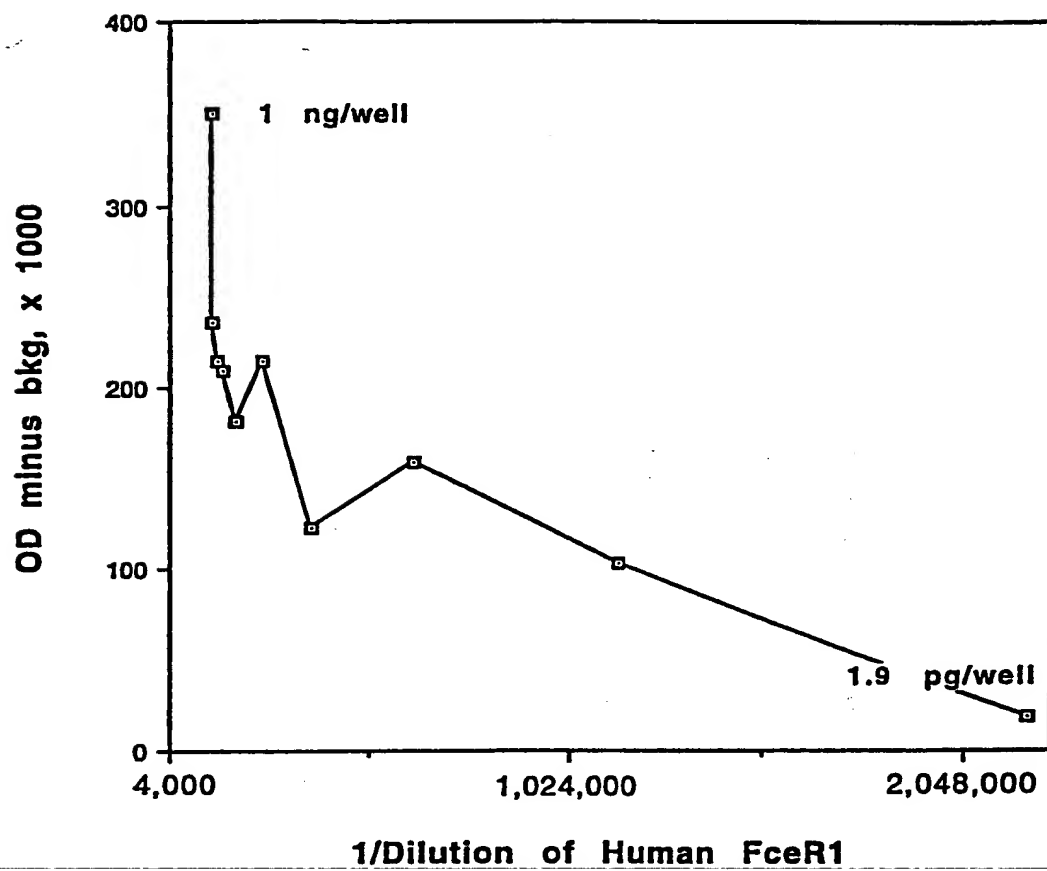


Fig. 4

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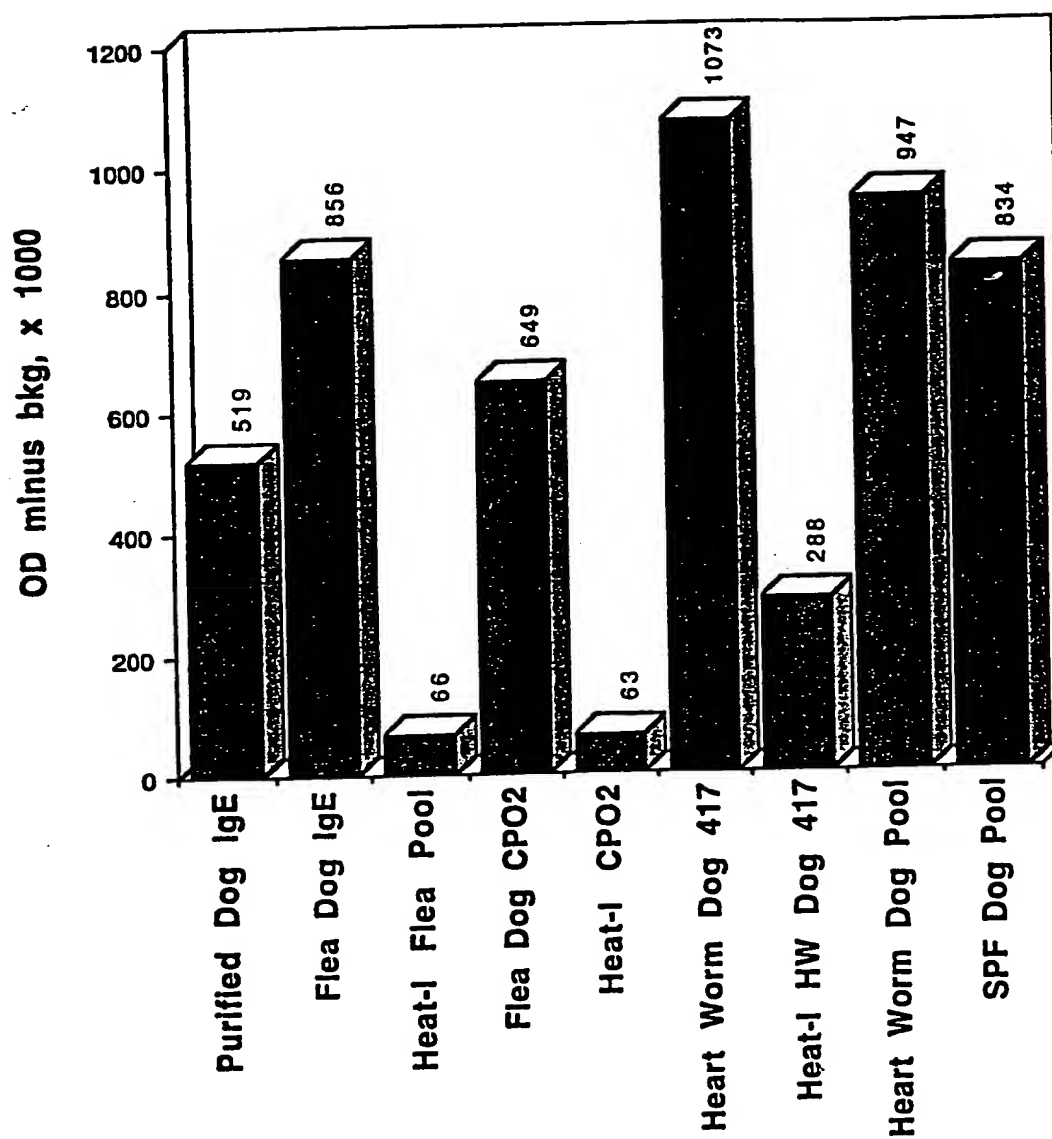


Fig. 5

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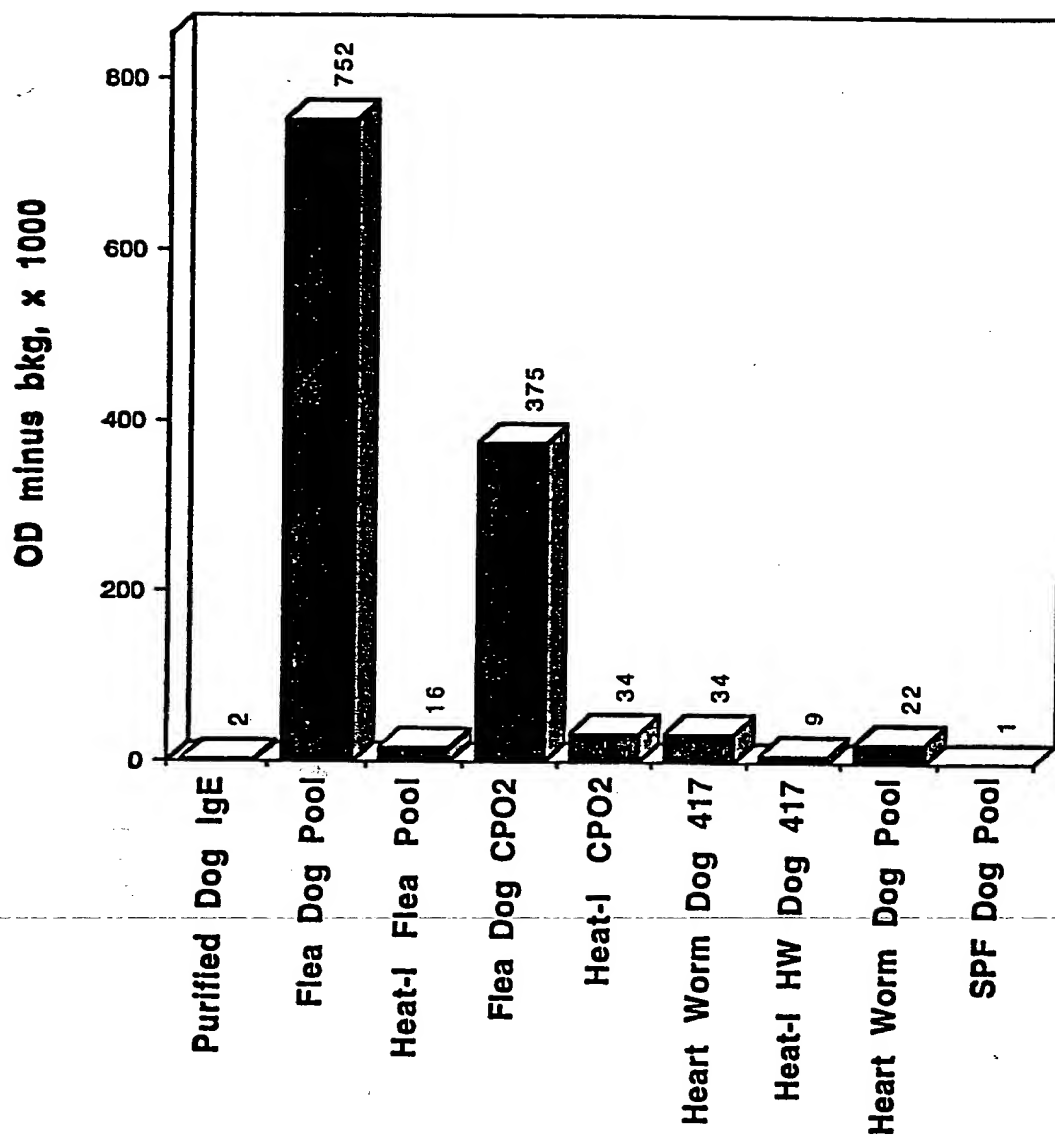


Fig. 6

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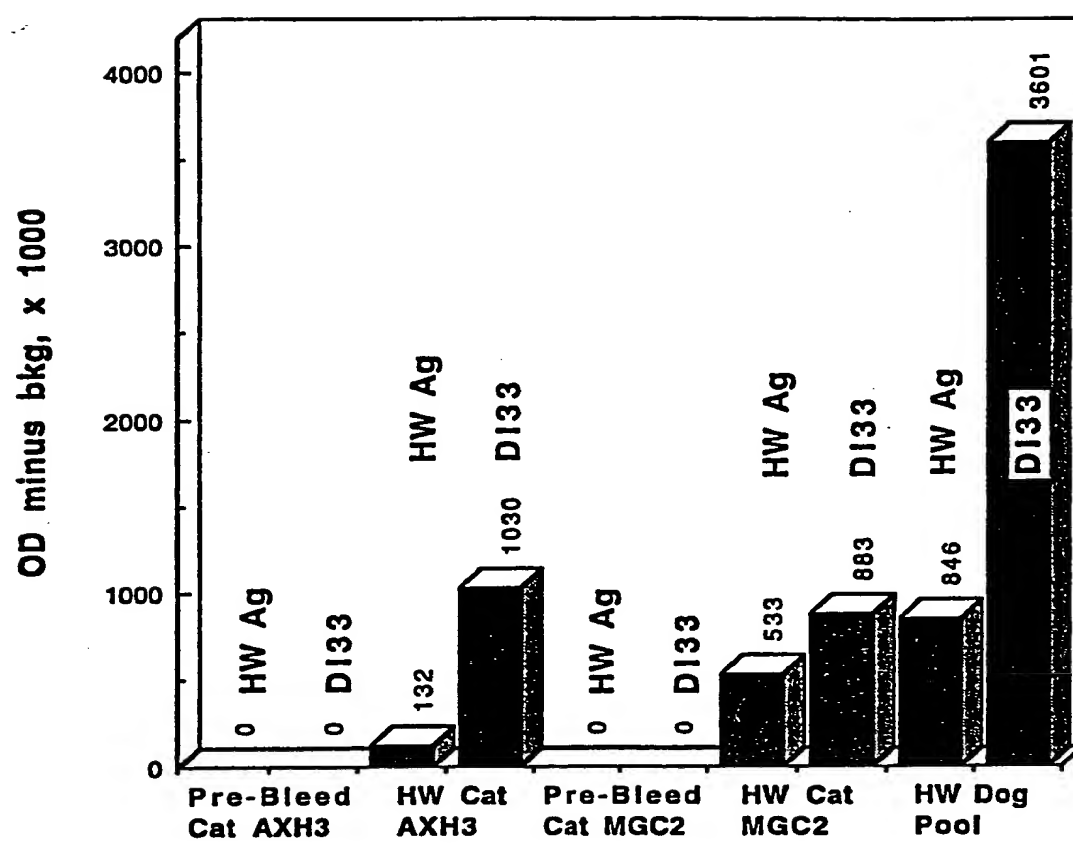


Fig. 7

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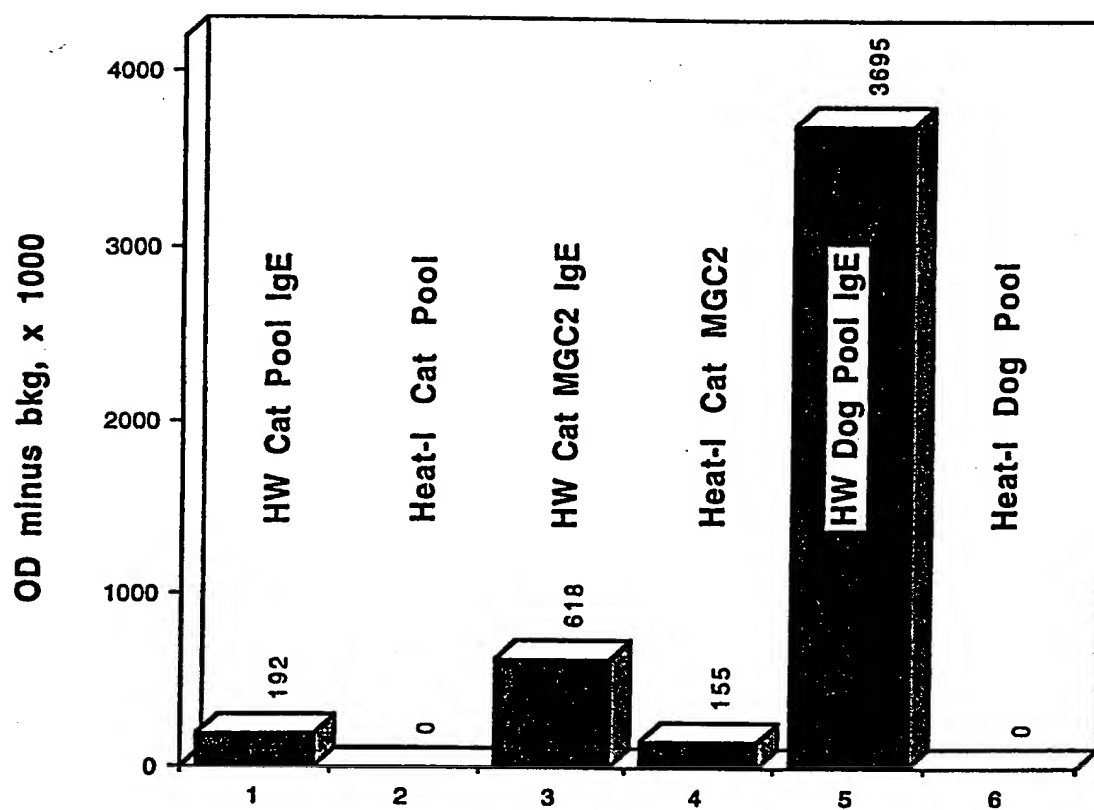


Fig. 8

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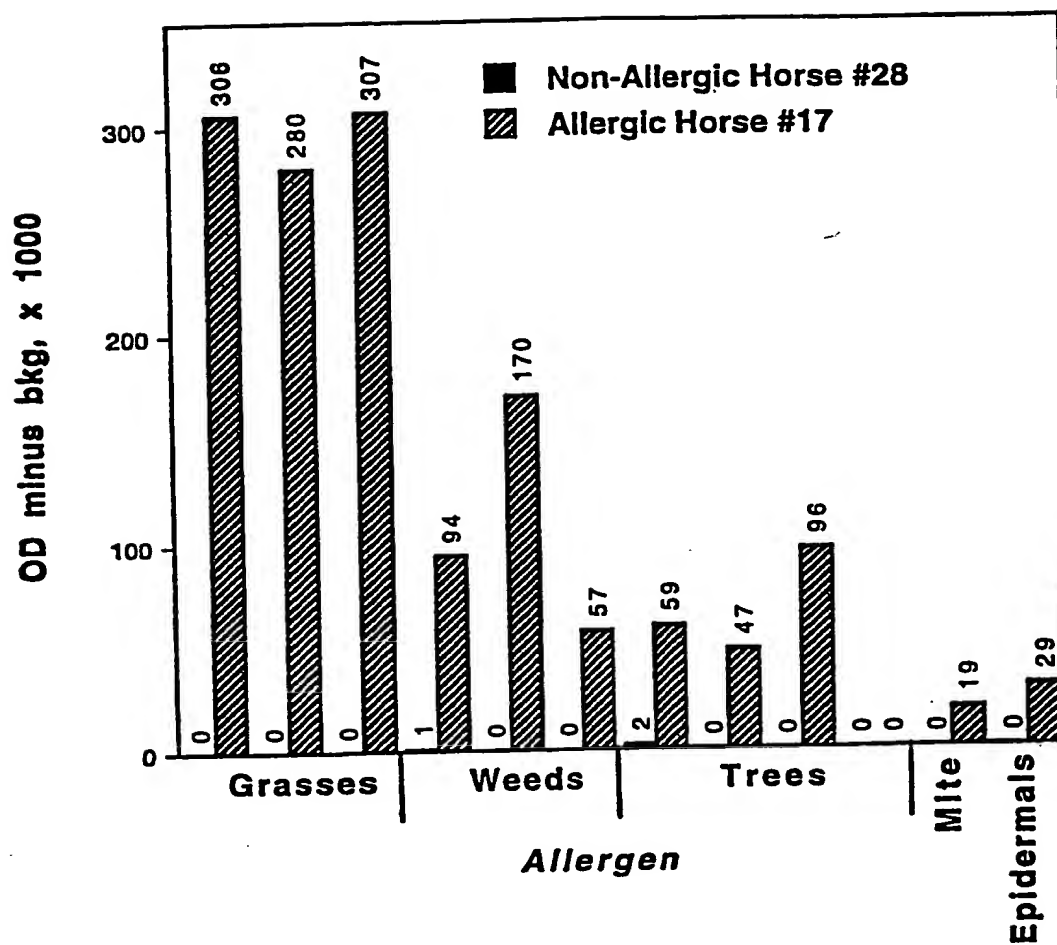
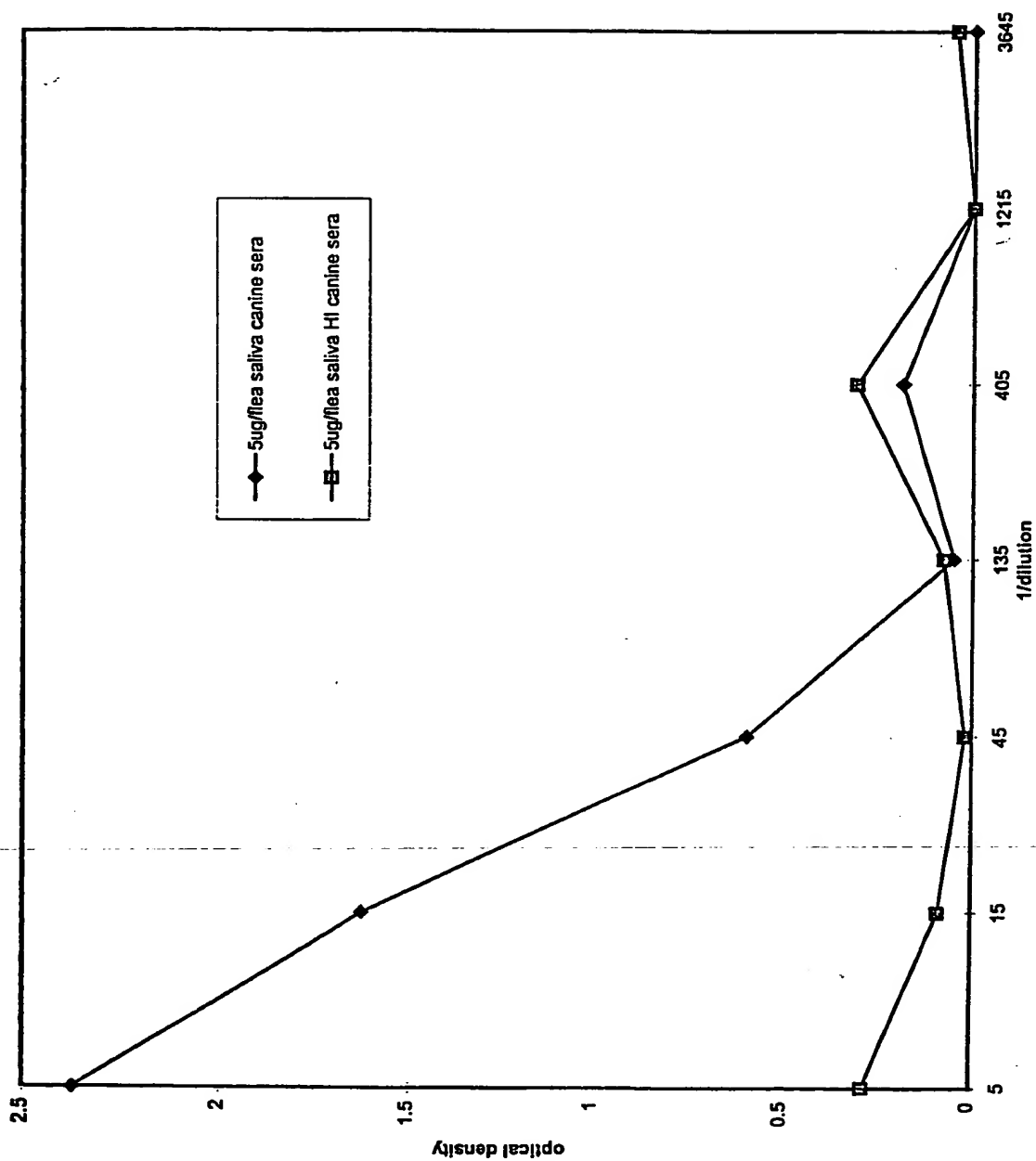


Fig. 9

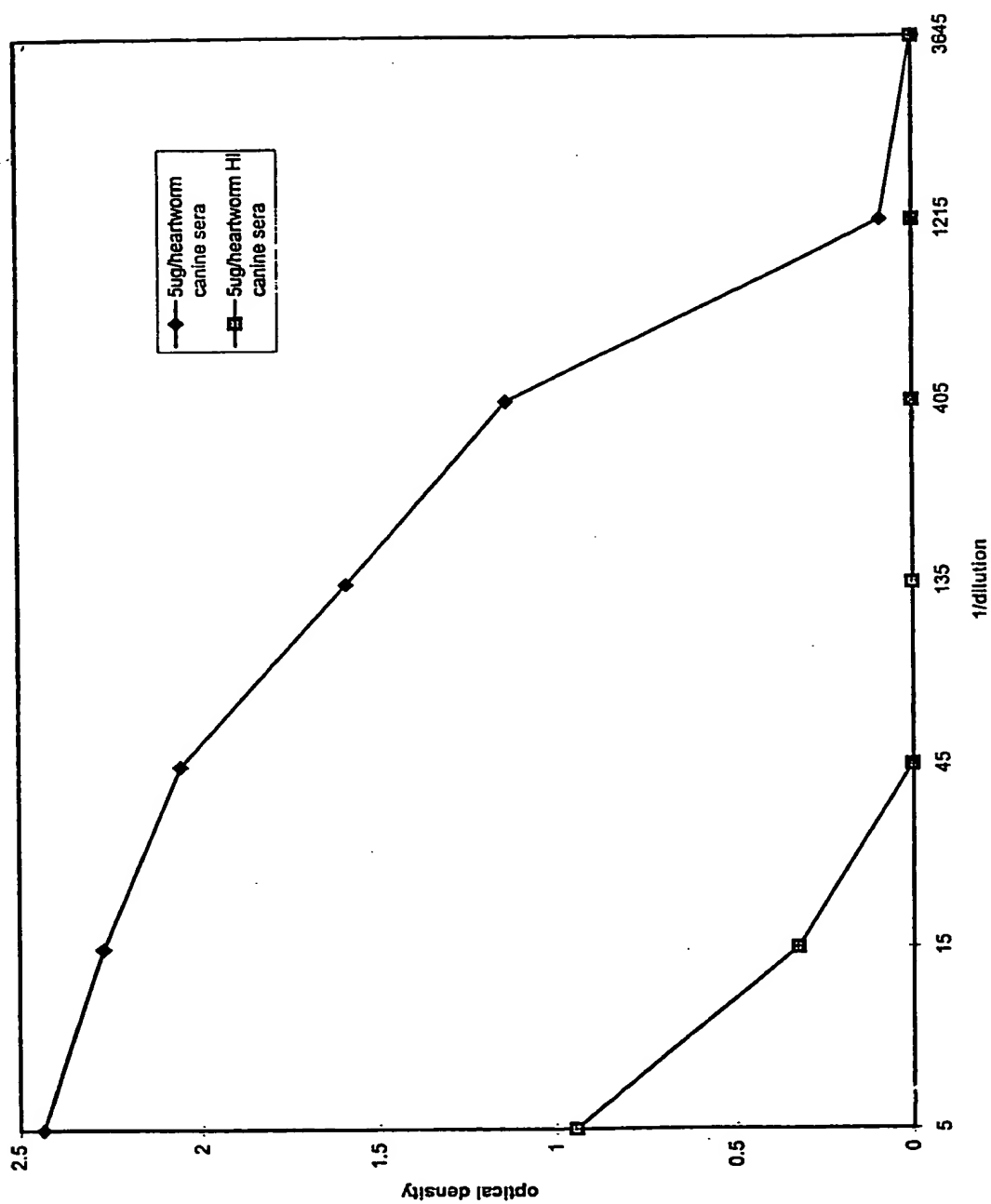
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Fig. 10



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Fig. 11



INTERNATIONAL SEARCH REPORT

Int. .ional Application No

PCT/US 97/21651

A. CLASSIFICATION OF SUBJECT MATTER

IPC 6 G01N33/68 G01N33/566

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 G01N C07K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P, X	WO 97 24617 A (NOVARTIS/SANDOZ) 10 July 1997 see claims see page 2, paragraph 2 see page 5, paragraph 2	1-69
X	PATENT ABSTRACTS OF JAPAN vol. 095, no. 007, 31 August 1995 & JP 07 092167 A (KINKI UNIV; OTHERS: 01), 7 April 1995, see abstract	1-69
X	PATENT ABSTRACTS OF JAPAN vol. 095, no. 006, 31 July 1995 & JP 07 072150 A (TONEN CORP; OTHERS: 01), 17 March 1995, see abstract	1-69

☐ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

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Date of the actual completion of the international search

12 March 1998

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19/03/1998

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Routledge, B

INTERNATIONAL SEARCH REPORT

information on patent family members

International Application No.

PCT/US 97/21651

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9724617 A	10-07-97	AU 1305897 A	28-07-97
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